PATENT COOPERATION TREATY

From the INTERNATIONAL BUREAU

PCT

NOTIFICATION OF ELECTION

(PCT Rule 61.2)

To:

United States Patent and Trademark Office
(Box PCT)
Crystal Plaza 2
Washington, DC 20231
ÉTATS-UNIS D'AMÉRIQUE

in its capacity as elected Office

Date of mailing (day/month/year) 03 December 1998 (03.12.98)	
International application No. PCT/GB98/01238	**Applicant's or agent's file reference** PHM 70228/WO
International filing date (day/month/year) 28 April 1998 (28.04.98)	**Priority date** (day/month/year) 30 April 1997 (30.04.97)
Applicant HEERY, David, Michael et al	

1. The designated Office is hereby notified of its election made:

[X] in the demand filed with the International Preliminary Examining Authority on:

02 November 1998 (02.11.98)

[ ] in a notice effecting later election filed with the International Bureau on:

2. The election [X] was

[ ] was not

made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).

The International Bureau of WIPO **34, chemin des Colombettes** **1211 Geneva 20, Switzerland**	Authorized officer Nicola Wolff
Facsimile No.: (41-22) 740.14.35	Telephone No.: (41-22) 338.83.38

Form PCT/IB/331 (July 1992)

2373745

BEST AVAILABLE COPY

PATENT COOPERATION TREATY

PCT

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference	**FOR FURTHER ACTION** see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.	
PHM 70228/WO		
International application No.	International filing date *(day/month/year)*	(Earliest) Priority Date *(day/month/year)*
PCT/GB 98/ 01238	28/04/1998	30/04/1997

Applicant

IMPERIAL CANCER RESEARCH TECHNOLOGY LTD et al.

This International Search Report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This International Search Report consists of a total of 3 sheets.

[X] It is also accompanied by a copy of each prior art document cited in this report.

1. [ ] **Certain claims were found unsearchable** (see Box I).

2. [ ] **Unity of invention is lacking** (see Box II).

3. [X] The international application contains disclosure of a **nucleotide and/or amino acid sequence listing** and the international search was carried out on the basis of the sequence listing
   - [X] filed with the international application.
   - [ ] furnished by the applicant separately from the international application,
     - [ ] but not accompanied by a statement to the effect that it did not include matter going beyond the disclosure in the international application as filed.
   - [ ] Transcribed by this Authority

4. With regard to the **title,**
   - [ ] the text is approved as submitted by the applicant
   - [X] the text has been established by this Authority to read as follows:

   INHIBITORS OF NUCLEAR PROTEIN: NUCLEAR RECEPTOR INTERACTION

5. With regard to the **abstract,**
   - [ ] the text is approved as submitted by the applicant
   - [X] the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this International Search Report, submit comments to this Authority.

6. The figure of the **drawings** to be published with the abstract is:

   Figure No. _____
   - [ ] as suggested by the applicant.
   - [ ] because the applicant failed to suggest a figure.
   - [ ] because this figure better characterizes the invention.

   [X] None of the figures.

Box III TEXT OF THE ABSTRACT (Continuation of item 5 of the first sheet)

- line 14, delete all remaining text after "repression of target genes" and replace semi-colon with full stop.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/GB 98/01238

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9811907 A	26-03-1998	AU 4345597 A	14-04-1998
WO 9802455 A	22-01-1998	NONE	
WO 9745737 A	04-12-1997	NONE	

INTERNATIONAL SEARCH REPORT

International Application No
PCT/GB 98/01238

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 G01N33/68 C07K7/06 C07K7/08

According to International Patent Classification(IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 G01N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	WO 98 11907 A (SQUIBB BRISTOL MYERS CO) 26 March 1998 see claims 1,7,14,15,21 see page 13, line 1 - line 12 see page 15, line 5 - page 25	1-19
P,X	WO 98 02455 A (INST NAT SANTE RECH MED ;UNIV PASTEUR (FR); CENTRE NAT RECH SCIENT) 22 January 1998 see claims 14-19 see page 34, line 4 - page 43, line 2	1-19
P,X	WO 97 45737 A (AMERICAN CYANAMID CO) 4 December 1997 see claims 1-16 see page 4, line 19 - line 26 see page 8, line 18 - page 9, line 7	1-19

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

° Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
12 August 1998	20/08/1998

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Routledge, B

1

PCT

REQUEST

The undersigned requests that the present international application be processed according to the Patent Cooperation Treaty.

For receiving Office use only

International Application No.

International Filing Date

Name of receiving Office and "PCT International Application"

Applicant's or agent's file reference *(if desired) (12 characters maximum)* PHM 70228/WO

Box No. I TITLE OF INVENTION

CHEMICAL COMPOUNDS

Box No. II APPLICANT

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (i.e. country) of residence if no State of residence is indicated below.)*

Imperial Cancer Research Technology Limited
Sardinia House
Sardinia Street
London
WC2A 3NL
GB

☐ This person is also inventor.

Telephone No. +44-171-269-3280

Facsimile No. +44-171-269-3094

Teleprinter No.

State *(i.e. country)* of nationality: GB

State *(i.e. country)* of residence: GB

This person is applicant for the purposes of: ☐ all designated States ☒ all designated States except the United States of America ☐ the United States of America only ☐ the States indicated in the Supplemental Box

Box No. III FURTHER APPLICANT(S) AND/OR (FURTHER) INVENTOR(S)

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (i.e. country) of residence if no State of residence is indicated below.)*

HEERY, David Michael
Molecular Endocrinology Laboratory
Imperial Cancer Research Fund
44 Lincoln's Inn Fields
London
WC2A 3PX, GB

This person is:

☐ applicant only

☒ applicant and inventor

☐ inventor only *(If this check-box is marked, do not fill in below.)*

State *(i.e. country)* of nationality: IE

State *(i.e. country)* of residence: GB

This person is applicant for the purposes of: ☐ all designated States ☐ all designated States except the United States of America ☒ the United States of America only ☐ the States indicated in the Supplemental Box

☒ Further applicants and/or (further) inventors are indicated on a continuation sheet.

Box No. IV AGENT OR COMMON REPRESENTATIVE; OR ADDRESS FOR CORRESPONDENCE

The person identified below is hereby/has been appointed to act on behalf of the applicant(s) before the competent International Authorities as: ☒ agent ☐ common representative

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country.)*

GILES, Allen Frank
Intellectual Property Department
ZENECA Pharmaceuticals
Mereside, Alderley Park
Macclesfield, Cheshire
SK10 4TG, GB

Telephone No. +44-1625-516753

Facsimile No. +44-1625-583358

Teleprinter No. 669095/669388 ZENPHA G

☐ Mark this check-box where no agent or common representative is/has been appointed and the space above is used instead to indicate a special address to which correspondence should be sent.

Continuation of Box No. III FURTHER APPLICANTS AND/OR (FURTHER) INVENTORS

If none of the following sub-boxes is used, this sheet is not to be included in the request.

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (i.e. country) of residence if no State of residence is indicated below.)*

PARKER, Malcolm George
Molecular Endocrinology Laboratory
Imperial Cancer Research Fund
44 Lincoln's Inn Fields
London
WC2A 3PX, GB

This person is:

- [ ] applicant only
- [x] applicant and inventor
- [ ] inventor only *(If this check-box is marked, do not fill in below.)*

State *(i.e. country)* of nationality: GB

State *(i.e. country)* of residence: GB

This person is applicant for the purposes of:
- [ ] all designated States
- [ ] all designated States except the United States of America
- [x] the United States of America only
- [ ] the States indicated in the Supplemental Box

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (i.e. country) of residence if no State of residence is indicated below.)*

This person is:

- [ ] applicant only
- [ ] applicant and inventor
- [ ] inventor only *(If this check-box is marked, do not fill in below.)*

State *(i.e. country)* of nationality:

State *(i.e. country)* of residence:

This person is applicant for the purposes of:
- [ ] all designated States
- [ ] all designated States except the United States of America
- [ ] the United States of America only
- [ ] the States indicated in the Supplemental Box

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (i.e. country) of residence if no State of residence is indicated below.)*

This person is:

- [ ] applicant only
- [ ] applicant and inventor
- [ ] inventor only *(If this check-box is marked, do not fill in below.)*

State *(i.e. country)* of nationality:

State *(i.e. country)* of residence:

This person is applicant for the purposes of:
- [ ] all designated States
- [ ] all designated States except the United States of America
- [ ] the United States of America only
- [ ] the States indicated in the Supplemental Box

Name and address: *(Family name followed by given name; for a legal entity, full official designation. The address must include postal code and name of country. The country of the address indicated in this Box is the applicant's State (i.e. country) of residence if no State of residence is indicated below.)*

This person is:

- [ ] applicant only
- [ ] applicant and inventor
- [ ] inventor only *(If this check-box is marked, do not fill in below.)*

State *(i.e. country)* of nationality:

State *(i.e. country)* of residence:

This person is applicant for the purposes of:
- [ ] all designated States
- [ ] all designated States except the United States of America
- [ ] the United States of America only
- [ ] the States indicated in the Supplemental Box

- [ ] Further applicants and/or (further) inventors are indicated on another continuation sheet.

Box No. V DESIGNATION OF STATES

The following designations are hereby made under Rule 4.9(a) *(mark the applicable check-boxes; at least one must be marked)*:

Regional Patent

- ☒ AP **ARIPO Patent:** GH Ghana, GM Gambia, KE Kenya, LS Lesotho, MW Malawi, SD Sudan, SZ Swaziland, UG Uganda, ZW Zimbabwe, and any other State which is a Contracting State of the Harare Protocol and of the PCT
- ☒ EA **Eurasian Patent:** AM Armenia, AZ Azerbaijan, BY Belarus, KG Kyrgyzstan, KZ Kazakhstan, MD Republic of Moldova, RU Russian Federation, TJ Tajikistan, TM Turkmenistan, and any other State which is a Contracting State of the Eurasian Patent Convention and of the PCT
- ☒ EP **European Patent:** AT Austria, BE Belgium, CH and LI Switzerland and Liechtenstein, DE Germany, DK Denmark, ES Spain, FI Finland, FR France, GB United Kingdom, GR Greece, IE Ireland, IT Italy, LU Luxembourg, MC Monaco, NL Netherlands, PT Portugal, SE Sweden, and any other State which is a Contracting State of the European Patent Convention and of the PCT
- ☒ OA **OAPI Patent:** BF Burkina Faso, BJ Benin, CF Central African Republic, CG Congo, CI Côte d'Ivoire, CM Cameroon, GA Gabon, GN Guinea, ML Mali, MR Mauritania, NE Niger, SN Senegal, TD Chad, TG Togo, and any other State which is a member State of OAPI and a Contracting State of the PCT *(if other kind of protection or treatment desired, specify on dotted line)* .....

National Patent *(if other kind of protection or treatment desired, specify on dotted line)*:

- ☒ AL Albania
- ☒ AM Armenia
- ☒ AT Austria
- ☒ AU Australia
- ☒ AZ Azerbaijan
- ☒ BA Bosnia and Herzegovina
- ☒ BB Barbados
- ☒ BG Bulgaria
- ☒ BR Brazil
- ☒ BY Belarus
- ☒ CA Canada
- ☒ CH and LI Switzerland and Liechtenstein
- ☒ CN China
- ☒ CU Cuba
- ☒ CZ Czech Republic
- ☒ DE Germany
- ☒ DK Denmark
- ☒ EE Estonia
- ☒ ES Spain
- ☒ FI Finland
- ☒ GB United Kingdom
- ☒ GE Georgia
- ☒ GH Ghana
- ☒ GM Gambia
- ☒ GW Guinea-Bissau
- ☒ HU Hungary
- ☒ ID Indonesia
- ☒ IL Israel
- ☒ IS Iceland
- ☒ JP Japan
- ☒ KE Kenya
- ☒ KG Kyrgyzstan
- ☒ KP Democratic People's Republic of Korea
- ☒ KR Republic of Korea
- ☒ KZ Kazakhstan
- ☒ LC Saint Lucia
- ☒ LK Sri Lanka
- ☒ LR Liberia
- ☒ LS Lesotho
- ☒ LT Lithuania
- ☒ LU Luxembourg
- ☒ LV Latvia
- ☒ MD Republic of Moldova
- ☒ MG Madagascar
- ☒ MK The former Yugoslav Republic of Macedonia
- ☒ MN Mongolia
- ☒ MW Malawi
- ☒ MX Mexico
- ☒ NO Norway
- ☒ NZ New Zealand
- ☒ PL Poland
- ☒ PT Portugal
- ☒ RO Romania
- ☒ RU Russian Federation
- ☒ SD Sudan
- ☒ SE Sweden
- ☒ SG Singapore
- ☒ SI Slovenia
- ☒ SK Slovakia
- ☒ SL Sierra Leone
- ☒ TJ Tajikistan
- ☒ TM Turkmenistan
- ☒ TR Turkey
- ☒ TT Trinidad and Tobago
- ☒ UA Ukraine
- ☒ UG Uganda
- ☒ US United States of America
- ☒ UZ Uzbekistan
- ☒ VN Viet Nam
- ☒ YU Yugoslavia
- ☒ ZW Zimbabwe

Check-boxes reserved for designating States (for the purposes of a national patent) which have become party to the PCT after issuance of this sheet:

- ☒ CY Cyprus
- ☐
- ☐

In addition to the designations made above, the applicant also makes under Rule 4.9(b) all designations which would be permitted under the PCT except the designation(s) of _____

The applicant declares that those additional designations are subject to confirmation and that any designation which is not confirmed before the expiration of 15 months from the priority date is to be regarded as withdrawn by the applicant at the expiration of that time limit. *(Confirmation of a designation consists of the filing of a notice specifying that designation and the payment of the designation and confirmation fees. Confirmation must reach the receiving Office within the 15-month time limit.)*

Box No. VI PRIORITY CLAIM Further priority claims are indicated in the Supplemental Box ☐

The priority of the following earlier application(s) is hereby claimed:

Country *(in which, or for which, the application was filed)*	Filing Date *(day/month/year)*	Application No.	Office of filing *(only for regional or international application)*
item (1) GB	30 April 1997 (30/04/97)	9708676.3	
item (2)			
item (3)			

Mark the following check-box if the certified copy of the earlier application is to be issued by the Office which for the purposes of the present international application is the receiving Office (a fee may be required):

☒ The receiving Office is hereby requested to prepare and transmit to the International Bureau a certified copy of the earlier application(s) identified above as item(s) : (1)

Box No. VII INTERNATIONAL SEARCHING AUTHORITY

Choice of International Searching Authority (ISA) *(If two or more International Searching Authorities are competent to carry out the international search, indicate the Authority chosen; the two-letter code may be used):* ISA /

Earlier search *Fill in where a search (international, international-type or other) by the International Searching Authority has already been carried out or requested and the Authority is now requested to base the international search, to the extent possible, on the results of that earlier search. Identify such search or request either by reference to the relevant application (or the translation thereof) or by reference to the search request:*

Country (or regional Office): Date *(day/month/year)*: Number:

Box No. VIII CHECK LIST

This international application contains the following number of sheets:

1. request	04	sheets
2. description	45	sheets
3. claims	03	sheets
4. abstract	01	sheets
5. drawings	08	sheets
Total	61	sheets

This international application is accompanied by the item(s) marked below:

1. ☒ separate signed power of attorney
2. ☐ copy of general power of attorney
3. ☐ statement explaining lack of signature
4. ☐ priority document(s) *identified in Box No. VI as item(s):*
5. ☒ fee calculation sheet
6. ☐ separate indications concerning deposited microorganisms
7. ☒ nucleotide and/or amino acid sequence listing (diskette)
8. ☐ other *(specify)*:

Figure No. _____ of the drawings (if any) should accompany the abstract when it is published.

Box No. IX SIGNATURE OF APPLICANT OR AGENT

Next to each signature, indicate the name of the person signing and the capacity in which the person signs (if such capacity is not obvious from reading the request).

A. F. Giles.

GILES, ALLEN FRANK
AUTHORISED REPRESENTATIVE

For receiving Office use only

1. Date of actual receipt of the purported international application:
2. Drawings: ☐ received: ☐ not received:
3. Corrected date of actual receipt due to later but timely received papers or drawings completing the purported international application:
4. Date of timely receipt of the required corrections under PCT Article 11(2):
5. International Searching Authority specified by the applicant: ISA /
6. ☐ Transmittal of search copy delayed until search fee is paid

For International Bureau use only

Date of receipt of the record copy by the International Bureau:

PATENT COOPERATION TREATY

17 AUG 1999

16 AUG 1999

From the
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

To:

GILES, A.
ZENECA Pharmaceuticals
Intellectual Property Department
Mereside, Alderley Park
Macclesfield
Cheshire SK10 4TG
GRANDE BRETAGNE

RECEIVED
12 AUG 1999
ZENECA PLC
INTELLECTUAL PROPERTY
GLUCE

PCT

NOTIFICATION OF TRANSMITTAL OF THE INTERNATIONAL PRELIMINARY EXAMINATION REPORT
(PCT Rule 71.1)

Date of mailing *(day/month/year)*	04.08.99
Applicant's or agent's file reference PHM 70228/WO	**IMPORTANT NOTIFICATION**

International application No.	International filing date *(day/month/year)*	Priority date *(day/month/year)*
PCT/GB98/01238	28/04/1998	30/04/1997

Applicant
IMPERIAL CANCER RESEARCH TECHNOLOGY LTD et al.

1. The applicant is hereby notified that this International Preliminary Examining Authority transmits herewith the international preliminary examination report and its annexes, if any, established on the international application.

2. A copy of the report and its annexes, if any, is being transmitted to the International Bureau for communication to all the elected Offices.

3. Where required by any of the elected Offices, the International Bureau will prepare an English translation of the report (but not of any annexes) and will transmit such translation to those Offices.

4. **REMINDER**

The applicant must enter the national phase before each elected Office by performing certain acts (filing translations and paying national fees) within 30 months from the priority date (or later in some Offices) (Article 39(1)) (see also the reminder sent by the International Bureau with Form PCT/IB/301).

Where a translation of the international application must be furnished to an elected Office, that translation must contain a translation of any annexes to the international preliminary examination report. It is the applicant's responsibility to prepare and furnish such translation directly to each elected Office concerned.

For further details on the applicable time limits and requirements of the elected Offices, see Volume II of the PCT Applicant's Guide.

Name and mailing address of the IPEA/	Authorized officer
European Patent Office - P.B. 5818 Patentlaan 2 NL-2280 HV Rijswijk - Pays Bas Tel. (+31-70) 340-2040 Tx: 31 651 epo nl Fax: (+31-70) 340-3016	Kruydenberg, G Tel.(+31-70)-340-2277

Form PCT/IPEA/416 (July 1992)

PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
PHM 70228/WO		
International application No.	International filing date *(day/month/year)*	Priority date *(day/month/year)*
PCT/GB98/01238	28/04/1998	30/04/1997
International Patent Classification (IPC) or national classification and IPC G01N33/68		
Applicant IMPERIAL CANCER RESEARCH TECHNOLOGY LTD et al.		

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.

2. This REPORT consists of a total of 5 sheets, including this cover sheet.

   ☐ This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

   These annexes consist of a total of sheets.

3. This report contains indications relating to the following items:

   - I ☒ Basis of the report
   - II ☐ Priority
   - III ☐ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
   - IV ☐ Lack of unity of invention
   - V ☒ Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations suporting such statement
   - VI ☒ Certain documents cited
   - VII ☒ Certain defects in the international application
   - VIII ☒ Certain observations on the international application

Date of submission of the demand	Date of completion of this report
02/11/1998	04.08.99
Name and mailing address of the international preliminary examining authority: European Patent Office - P.B. 5818 Patentlaan 2 NL-2280 HV Rijswijk - Pays Bas Tel. (+31-70) 340-2040 Tx: 31 651 epo nl Fax: (+31-70) 340-3016	Authorized officer Routledge, B Telephone No. (+31-70)-340

Form PCT/IPEA/409 (cover sheet) (January 1994)

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/GB98/01238

I. Basis of the report

1. This report has been drawn on the basis of (*substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments.*):

Description, pages:

1-45 as originally filed

Claims, No.:

1-19 as originally filed

Drawings, sheets:

1/8-8/8 as originally filed

2. The amendments have resulted in the cancellation of:

☐ the description, pages:
☐ the claims, Nos.:
☐ the drawings, sheets:

3. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

4. Additional observations, if necessary:

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes:	Claims	1-19
	No:	Claims	
Inventive step (IS)	Yes:	Claims	1-19
	No:	Claims	
Industrial applicability (IA)	Yes:	Claims	1-19
	No:	Claims	

2. Citations and explanations

see separate sheet

VI. Certain documents cited

1. Certain published documents (Rule 70.10)

and / or

2. Non-written disclosures (Rule 70.9)

see separate sheet

VII. Certain defects in the international application

The following defects in the form or contents of the international application have been noted:

see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

see separate sheet

Re Item V

Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. The application meets the criteria of Article 33(2) and (3) PCT in that a method for identifying inhibitors of the reaction between a nuclear receptor (NR) or fragment thereof, such as a transcription factor which binds to a nuclear protein (NP) or fragment thereof via a signature motif on the NP, the binding between the NP and the NR being part of activation or repression of a target gene and the NP being a bridging factor between the NR and a transcription initiation complex, is neither taught nor suggested in the cited prior art.

2. The application meets the criteria of Article 33(3) PCT with respect to industrial applicability.

Re Item VI

Certain documents cited

Certain published documents (Rule 70.10)

Application No Patent No	Publication date *(day/month/year)*	Filing date *(day/month/year)*	Priority date *(valid claim)* *(day/month/year)*
WO 98/11907	**26.03.98**	**12.09.97**	**20.09.96**
WO 98/02455	**22.01.98**	**11.07.97**	**12.07.96**
WO 97/45737	**04.12.97**	**30.05.97**	**31.05.96**

1. **WO 98/11907** discloses peptide inhibitors of nuclear protein translocation having localization sequences, **WO 98/02455** discloses a screening method for identifying agonists and antagonists of the AD1 activation domain activity of TIF2 and **WO 97/45737** discloses the screening of inhibitors of orphan nuclear receptors, especially the insect ultraspiracle protein.

Re Item VII

Certain defects in the international application

1. The application does not meet the requirements of Rule 11.13(m) PCT with regard to the labelling of the wild type in Figure **2A** and the equivalent passage at line **22** page.

Re Item VIII

Certain observations on the international application

1. The application does met the criteria of Article 6 PCT for the following reasons:-

   (a) The use of the relative term "short" in claims **1** and **14** (and description) leads to a lack of clarity.

   (b) Claim **13** lacks clarity as a "potential inhibitor" may be derived from a "peptide library" but may not itself be a library, as an inhibitor is a single compound.

   (c) Claim **19** lacks clarity as the claim refers to signature motifs "..as defined in any one of claims **1-6**..". However, claim **1** does not contain a defined signature motif.

   (d) The use of the phrase "..incorporated herein by reference.." page **9** leads to a lack of clarity.

2. The unit **MICRON** on page **24** is not expressed in terms of the units stipulated by Rule 10.1/(a)/and/(b) PCT. In the SI system the unit is usually written "µm" or "micrometres".

3. The term **TWEEN** on page **21** appears to be a registered trade mark but has not been acknowledged as such.

19

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INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
PHM 70228/WO		
International application No.	International filing date *(day/month/year)*	Priority date *(day/month/year)*
PCT/GB98/01238	28/04/1998	30/04/1997
International Patent Classification (IPC) or national classification and IPC G01N33/68		
Applicant IMPERIAL CANCER RESEARCH TECHNOLOGY LTD et al.		

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.

2. This REPORT consists of a total of 5 sheets, including this cover sheet.

   ☐ This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

   These annexes consist of a total of sheets.

3. This report contains indications relating to the following items:

   - I ☒ Basis of the report
   - II ☐ Priority
   - III ☐ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
   - IV ☐ Lack of unity of invention
   - V ☒ Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations suporting such statement
   - VI ☒ Certain documents cited
   - VII ☒ Certain defects in the international application
   - VIII ☒ Certain observations on the international application

Date of submission of the demand	Date of completion of this report
02/11/1998	04.08.99
Name and mailing address of the international preliminary examining authority: European Patent Office - P.B. 5818 Patentlaan 2 NL-2280 HV Rijswijk - Pays Bas Tel. (+31-70) 340-2040 Tx: 31 651 epo nl Fax: (+31-70) 340-3016	Authorized officer Routledge, B Telephone No. (+31-70)-340

Form PCT/IPEA/409 (cover sheet) (January 1994)

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/GB98/01238

I. Basis of the report

1. This report has been drawn on the basis of (*substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments.*):

Description, pages:

1-45 as originally filed

Claims, No.:

1-19 as originally filed

Drawings, sheets:

1/8-8/8 as originally filed

2. The amendments have resulted in the cancellation of:

☐ the description, pages:
☐ the claims, Nos.:
☐ the drawings, sheets:

3. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

4. Additional observations, if necessary:

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes:	Claims	1-19
	No:	Claims	
Inventive step (IS)	Yes:	Claims	1-19
	No:	Claims	
Industrial applicability (IA)	Yes:	Claims	1-19
	No:	Claims	

2. Citations and explanations

see separate sheet

VI. Certain documents cited

1. Certain published documents (Rule 70.10)

and / or

2. Non-written disclosures (Rule 70.9)

see separate sheet

VII. Certain defects in the international application

The following defects in the form or contents of the international application have been noted:

see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

see separate sheet

Re Item V

Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. The application meets the criteria of Article 33(2) and (3) PCT in that a method for identifying inhibitors of the reaction between a nuclear receptor (NR) or fragment thereof, such as a transcription factor which binds to a nuclear protein (NP) or fragment thereof via a signature motif on the NP, the binding between the NP and the NR being part of activation or repression of a target gene and the NP being a bridging factor between the NR and a transcription initiation complex, is neither taught nor suggested in the cited prior art.

2. The application meets the criteria of Article 33(3) PCT with respect to industrial applicability.

Re Item VI

Certain documents cited

Certain published documents (Rule 70.10)

Application No Patent No	Publication date *(day/month/year)*	Filing date *(day/month/year)*	Priority date *(valid claim)* *(day/month/year)*
WO 98/11907	**26.03.98**	**12.09.97**	**20.09.96**
WO 98/02455	**22.01.98**	**11.07.97**	**12.07.96**
WO 97/45737	**04.12.97**	**30.05.97**	**31.05.96**

1. **WO 98/11907** discloses peptide inhibitors of nuclear protein translocation having localization sequences, **WO 98/02455** discloses a screening method for identifying agonists and antagonists of the AD1 activation domain activity of TIF2 and **WO 97/45737** discloses the screening of inhibitors of orphan nuclear receptors, especially the insect ultraspiracle protein.

Re Item VII

Certain defects in the international application

1. The application does not meet the requirements of Rule 11.13(m) PCT with regard to the labelling of the wild type in Figure **2A** and the equivalent passage at line **22** page.

Re Item VIII

Certain observations on the international application

1. The application does met the criteria of Article 6 PCT for the following reasons:-

   (a) The use of the relative term "short" in claims **1** and **14** (and description) leads to a lack of clarity.

   (b) Claim **13** lacks clarity as a "potential inhibitor" may be derived from a "peptide library" but may not itself be a library, as an inhibitor is a single compound.

   (c) Claim **19** lacks clarity as the claim refers to signature motifs "..as defined in any one of claims **1-6**..". However, claim **1** does not contain a defined signature motif.

   (d) The use of the phrase "..incorporated herein by reference.." page **9** leads to a lack of clarity.

2. The unit **MICRON** on page **24** is not expressed in terms of the units stipulated by Rule 10.1/(a)/and/(b) PCT. In the SI system the unit is usually written "µm" or "micrometres".

3. The term **TWEEN** on page **21** appears to be a registered trade mark but has not been acknowledged as such.

INTERNATIONAL SEARCH REPORT

Int. .ational Application No
PCT/GB 98/01238

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 G01N33/68 C07K7/06 C07K7/08

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 G01N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	WO 98 11907 A (SQUIBB BRISTOL MYERS CO) 26 March 1998 see claims 1,7,14,15,21 see page 13, line 1 - line 12 see page 15, line 5 - page 25	1-19
P,X	WO 98 02455 A (INST NAT SANTE RECH MED ;UNIV PASTEUR (FR); CENTRE NAT RECH SCIENT) 22 January 1998 see claims 14-19 see page 34, line 4 - page 43, line 2	1-19
P,X	WO 97 45737 A (AMERICAN CYANAMID CO) 4 December 1997 see claims 1-16 see page 4, line 19 - line 26 see page 8, line 18 - page 9, line 7	1-19

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

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"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
12 August 1998	20/08/1998

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Routledge, B

1

Form PCT/ISA/210 (second sheet) (July 1992)

INTERNATIONAL SEARCH REPORT

Information on patent family members

Int. donal Application No
PCT/GB 98/01238

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9811907 A	26-03-1998	AU 4345597 A	14-04-1998
WO 9802455 A	22-01-1998	NONE	
WO 9745737 A	04-12-1997	NONE	

PCT WORLD INTELLECTUAL PROPERTY ORGANIZATION
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(51) International Patent Classification 6 : G01N 33/68, C07K 7/06, 7/08

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(11) International Publication Number: **WO 98/49561**

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(30) Priority Data:
9708676.3 30 April 1997 (30.04.97) GB

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(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).

Published
With international search report.

(54) Title: INHIBITORS OF NUCLEAR PROTEIN/NUCLEAR RECEPTOR INTERACTION

(57) Abstract

A method for identifying inhibitor compounds capable of reducing the interaction between a first region which is a signature motif on a nuclear protein, and a second region which is that part of a nuclear receptor which is capable of interacting with the nuclear protein through binding to the signature motif, wherein: the nuclear protein is a bridging factor that is responsible for the interaction between a liganded nuclear receptor and a transcription initiation complex involved in regulation of gene expression; the nuclear receptor is a transcription factor, the signature motif is a short sequence of amino acid residues which is the key structural element of a nuclear protein which binds to a liganded nuclear receptor as part of the process of the activation or repression of target genes.

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INHIBITORS OF NUCLEAR PROTEIN: NUCLEAR RECEPTOR INTERACTION

The present invention relates to inhibition of the interaction between nuclear proteins and nuclear receptors through identification of the key structural element responsible for the interaction.

The binding of lipophilic hormones, retinoids and vitamins to members of the nuclear receptor (NR) superfamily (to form "liganded" receptors) modifies their DNA binding and transcriptional properties, resulting in the activation or repression of target genes [1,2]. Ligand binding induces conformational changes in NRs and promotes their association with a diverse group of nuclear proteins, including SRC-1/p160 [3,4,5], TIF2 [6,7] and CBP/p300 [4,5,8,9] which function as coactivators, and RIP-140 [10], TIF1 [11] and TRIP1/SUG1 [12, 13] whose functions are unclear.

The recruitment of nuclear proteins (coactivators and/or other so-called bridging proteins) by NRs is thought to be essential to their function as ligand-induced transcription factors. Structural studies of the ligand binding domains (LBDs) of three different nuclear hormone receptors, the retinoid X receptor α (RXRα) [15], the retinoic acid receptor γ (RARγ) [16] and the thyroid hormone receptor β (TRβ) [17], have led to the proposal that binding of ligand results in a realignment of a conserved amphipathic α-helix, Helix 12 (H12), generating a novel surface required for coactivator binding and consequently activator function 2 (AF2)-dependent transactivation. Consistent with this, mutations of conserved hydrophobic residues in H12 which impair AF2 [14, 18-20], also interfere with the ability of NRs to bind coactivators [4,6,10,11,13]. Less is known about the coactivator sequences which mediate interaction with NRs although several proteins appear to contain multiple NR binding sites [5,8,21]. Le Douarin *et al* (1996) in EMBO Journal, 15, 6701-6715, identified a leucine rich region in three coactivators (TIF1, RIP140 & TRIP3) which they called the "NR box"; see Figure 3D therein. However the present state of knowledge is completely silent about precisely how liganded nuclear receptors interact with nuclear proteins as a class to modify their DNA binding and transcriptional properties, resulting in the activation or repression of target genes. Indeed a commentator on the field stated, after the first filing date of the present invention, that "*characterizing the mechanisms by which nuclear factors engage the transcriptional apparatus in response to hormonal stimulation has seemed, at times, to be an*

insurmountable task" (Marc Montminy in Nature, 12th June, 1997, 387, 654-655, see 1st paragraph thereof).

The present invention is based on the discovery that a short signature motif present in the nuclear proteins is necessary and sufficient to mediate their binding to liganded NRs.

According to one aspect of the present invention there is provided a method for identifying inhibitor compounds capable of reducing the interaction between:

a) a first region which is a signature motif on a nuclear protein, and

b) a second region which is that part of a nuclear receptor which is capable of interacting with the nuclear protein through binding to the signature motif,

wherein:

the nuclear protein is a bridging factor that is responsible for the interaction between a liganded nuclear receptor and a transcription initiation complex involved in regulation of gene expression;

the nuclear receptor is a transcription factor;

the signature motif is a short sequence of amino acid residues which is the key structural element of a nuclear protein which binds to a liganded nuclear receptor as part of the process of the activation or repression of target genes; and

in which the method comprises taking:

i) the potential inhibitor compound;

ii) the liganded nuclear receptor or a fragment thereof in which the fragment comprises the second region defined in this claim in b) above;

iii) a nuclear protein fragment comprising a signature motif of the nuclear protein; and

iv) detecting the presence or absence of inhibition of the interaction between ii) and iii).

The term "nuclear protein" means the bridging factors (including coactivators) that are responsible for the interaction between a liganded nuclear receptor and the transcription initiation complex involved in regulation of gene expression (reviewed for steroid hormone receptors in Beato, M., Herrlich, P. & Schutz, G. *Cell* **83**, 851-857 (1995)). The term bridging factor may include part of the transcription initiation complex itself. The term "nuclear receptor" means the family of nuclear receptors such as described in Mangelsdorf, D.J., *et al. Cell* **83**, 835-839 (1995). The term "signature motif" means a short sequence of generally at least about 5 amino acids, preferably 4-10, more preferably 5-10 amino acid

residues which is the key structural element of a nuclear protein which binds to a liganded nuclear receptor as part of the process of the activation or repression of target genes. The term "liganded nuclear receptor" means an activated nuclear receptor, for example, with ligand bound thereto. The ligand can take various forms such as for example a hormone, a small molecule compound or a peptide. For example, in the case of some of the nuclear receptors (e.g. PPAR) there are non hormonal peptide ligands e.g. leukotrienes. Although nuclear receptors are generally activated through ligand binding, some receptors, such as for example orphan receptors, may be active without the need for ligand and/ or activated through non-ligand dependent pathways and these receptors are also within the scope of the invention as a less preferred embodiment.

The term "fragment" means an incomplete part. Before the present invention, a skilled person could not have known which fragment or fragments of a nuclear protein could be taken to retain activity. Use of fragments compared with whole proteins is particularly advantageous in screening assays. In a preferred embodiment a preferred fragment size of a nuclear protein is 8-10 amino acids, such as, for example, shown in Figure 1A herein. The liganded nuclear receptor is preferably in the form of a fragment. In general, fragments comprise at least 8 amino acids.

Preferably the signature motif is represented by B^1XXLL in which B^1 is any natural hydrophobic amino acid, L is leucine and X represents any natural amino acid. Values for X within the signature motif are independently selected i.e. X may be the same or different. Preferably B^1 is leucine or valine with leucine being most preferred. In some instances the preferred signature motif is further defined as B^2B^1XXLL wherein "B^2" is a hydrophobic amino acid residue as defined for B^1. A "natural hydrophobic amino acid" is defined as any one of isoleucine, leucine, methionine, phenylalanine, tryptophan, tyrosine or valine. Preferably the signature motif is in the conformation of a helix, preferably an amphipathic helix, and the leucine residues form a hydrophobic face thereof. Preferably the signature motif is positioned within a molecule so that it is available at the surface thereof for interaction with proteins. Preferably values of X do not include Cys or Pro. Preferably at least one value of X is not a natural hydrophobic amino acid or proline. One value of X is preferably independently selected from Arg, Asn, Asp, Glu, Gln, His, Lys, Ser, Thr, Gly or Ala, more preferably one value of X is independently selected from Arg, Asn, Asp, Glu, Gln,

His or Lys. Without wishing to be bound by theoretical considerations, it is believed that preferred values of X favour the signature motif forming an amphipathic helix.

After the first filing date of the present invention there was a simultaneous publication of the identification of LXXLL motifs by two groups (one of which included the inventors of the present invention), namely Heery *et al*, Nature, 12th June 1997, 387, 733-736 & Torchia *et al*, Nature, 12th June 1997, 387, 677-684.

Herein we show that the ability of a nuclear protein (SRC1) to bind a nuclear receptor (liganded ER) and enhance its transcriptional activity is dependent upon the integrity within the nuclear protein of the signature motif (LXXLL; SEQ ID NO: 1), as well as key hydrophobic residues in the conserved helix (Helix 12) of NRs required for their ligand-induced activation function (AF-2) [14]. The signature motif is also found in TIF1, TIF2, p300, RIP 140 and the TRIP proteins, and occurs within regions of these proteins known to be sufficient for interaction with NRs. Thus the LXXLL motif (SEQ ID NO: 1) is a signature sequence which facilitates the interaction of diverse proteins with nuclear receptors, and thus is a key part of a new family of nuclear proteins.

A preferred nuclear protein is a coactivator, in particular the nuclear protein includes any one of RIP 140, SRC-1, TIF2, CBP, p300, TIF1, Trip1, Trip2, Trip3, Trip4, Trip5, Trip8 or Trip9. Further preferred nuclear proteins include p/CIP, ARA70 & Trip230.

In this specification a reference to a nuclear protein or nuclear receptor includes isoforms thereof unless stated or otherwise implicit from the context. An isoform is one of a family or collection of related proteins derived from a single gene. Thus isoforms may differ slightly in their amino acid sequences such as for example from differential splicing of exons following transcription. SRC1a is an example of an isoform of SRC1. Two isoforms of SRC-1, namely SRC1a and SRC1e, have been shown to contain differences in number of signature motifs and to be functionally distinct in so far as they appear to play different roles in ER-mediated transcription (Kalkhoven *et al*, 1998, EMBO Journal, 17, 232-243).

Nuclear receptors are transcription factors. A preferred transcription factor comprises at least part of a conserved amphipathic α-helix, and especially preferred is retinoic acid receptor or a steroid hormone receptor. Preferred steroid hormone receptors are oestrogen receptor, progesterone receptor, androgen receptor and glucocorticoid receptor with oestrogen receptor being especially preferred.

Preferably the second region comprises at least part of a conserved amphipathic α-helix such as for example Helix 12 in the oestrogen receptor which is especially preferred.

An especially preferred combination of nuclear receptor and nuclear protein is one in which the nuclear receptor is oestrogen receptor and the nuclear protein is selected from SRC1, TIF2, CBP and p300, with SRC1 and especially SRC1a being most preferred.

A preferred method is in the form of a 2-hybrid assay system. Such assay systems are well known in the art; suitable references include Fields & Sternglanz (1994) TIG, August 1994, 10, 286-292 and US patent 5283173.

Any suitable assay design may be employed such as for example radioisotopic assay, scintillation proximity assay (reviewed by ND Cook, 1996, Drug Discovery Today, **1**, 287-294) or fluorescence, particularly time resolved fluorescence, assay (reviewed by MV Rogers, 1997, Drug Discovery Today, **2**, 156-160). High-throughput screeening technologies have been reviewed by Houston & Banks in Current Opinion in Biotechnology 1997, **8**, 734-740.

In a preferred embodiment of the invention the potential inhibitor is in the form of a peptide library based on a signature motif.

Encoded peptide libraries generally have a maximum of 20 possible amino acids at any one position. In practice, using current technology, it is difficult to screen libraries of more than 10^7- 10^8 members, which means that it is difficult to randomise more than 6 positions in a peptide. Hence, narrowing down the nuclear protein binding region to a signal motif is of great advantage if a peptide library approach is to be employed.

A peptide library is a collection of peptides of varying sequences. There are in general two ways to generate peptide libraries (reviewed by Scott, 1992; Birnbaum and Mosbach, 1992; Houghten, 1993;see also Abelson, 1996). The first approach is to generate libraries in which positive peptides are identified through the sequencing of the peptides themselves. Mixtures of peptides may be chemically synthesised in such a way that the peptides are linked to beads, so that each bead contains only one peptide. If a bead is identified which contains a positive peptide, the bead may be recovered and the peptide identified by chemical sequencing. This approach was first demonstrated using the ability of antibodies to identify specific six amino acid peptides from mixtures (Lam *et al*, 1991). The importance of using beads is that the identification event (in this case the antibody:peptide interaction) leads to the recovery of a bead which contains more peptide than that bound by the antibody itself. In a

related approach mixtures of free peptides can be synthesised and screened in pools; by using a deconvolution process, positive peptides are identified (Houghten *et al*, 1991).

The second approach is to generate libraries in which positive peptides are identified by sequencing a molecule which is associated in some way with the peptide. Peptides and some other molecule, such as a nucleic acid, can be cosynthesised on beads, so that each nucleic acid "tags" the peptide found on the same bead. If a bead is identified which is positive, sequencing the nucleic acid will identify the peptide on this bead (Brenner and Lerner, 1992). Alternatively, peptide libraries can be generated using the gene expression machinery of living organisms. In this approach, it is not necessary to make a library of peptide molecules. Instead a library of DNA molecules is constructed, so that each molecule encodes a peptide with a different sequence. This encoded library must then be expressed in a suitable host organism in order that the peptide library be produced. The library is then screened. It is essential that the nucleic acid which encodes the library remain physically linked to the protein in some way, so that recovery, or identification, of the active peptides leads to recovery of the DNA which encodes these peptides. The sequences of the active peptides may then be deduced by sequencing of the DNA which encodes them. Several variations of this approach have been described.

The mostly widely used version of this approach is to express peptides as part of the coat proteins of a virus such as M13. The viruses can be screened by the ability of this coat protein to bind target proteins (such as antibodies (Devlin *et al*, 1990; Scott and Smith, 1990; Cwirla *et al*, 1990) or receptors (the atrial natriuretic peptide receptor: Cunningham *et al* (1994) and the thrombopoietin receptor (Cwirla *et al*, 1997). The approach may also be used to find protease inhibitors through their ability to bind to proteases (Roberts *et al*, 1992; Markland *et al*, 1996) as well as to find optimal substrates for proteases, such as stromelysin and matrilysin (Smith *et al*, 1995) and subtilisin (Matthews and Wells, 1993)

An intracellular approach to the generation of peptides that recognise certain proteins is to use the yeast two-hybrid system. In the two-hybrid system, interacting proteins are fused to domains of transcription factors. If a protein:protein interaction occurs, then transcription of a reporter gene is stimulated (Fields and Song, 1989). By making one component of the two hybrid system a peptide library, and selecting for cells in which reporter gene output occurs, it is possible to isolate peptides which bind to a target protein. this approach was used to

identify peptides which bound to the retinoblastoma protein (Yang *et al*, 1995). In a similar approach, Colas *et al* (1996) expressed the peptide library as a loop in the surface of the E. coli TrxA protein and isolated peptides which bound to cyclin-dependent kinase 2 (Cdk2).

According to another aspect of the present invention there is provided a method of reducing the interaction between

a) a first region which is a signature motif on a nuclear protein, and

b) a second region which is that part of a nuclear receptor which is capable of interacting with the nuclear protein through binding to the signature motif,

in which the method comprises adding an inhibitor in the presence of the nuclear receptor and the nuclear protein, the inhibitor being characterised in that it reduces the interaction between the first region of the nuclear protein and the second region of the nuclear receptor.

According to another aspect of the present invention there is provided a novel inhibitor as described above. Preferably the inhibitor is a peptide, more preferably a peptide comprising the signature motif defined above, and more preferably the peptide has less than 15 amino acid residues. Especially preferred inhibitors are any one of the following peptides; PQAQQKSLLQQLLT (SEQ ID NO: 2), KLVQLLTTT (SEQ ID NO: 3), ILHRLLQE (SEQ ID NO: 4), or LLQQLLTE (SEQ ID NO: 5). Peptides may be prepared using conventional techniques for example using solid phase synthesis and Fmoc chemistry. These peptides are expected to be useful in the treatment of oestrogen responsive tumours. Inhibitors of the invention are expected to be useful in the treatment of any disease mediated through interaction between a signature motif on a nuclear protein and a nuclear receptor. For example, suitable inhibitors are expected to be useful in treatment of cancer or inflammation.

A novel inhibitor, for example, could be an antibody against a signature motif or a novel small molecule which binds to the signature motif or its complementary binding target (nuclear receptor second region) such that normal biological activity is prevented. Examples of small molecules include but are not limited to small peptides or peptide-like molecules.

Whilst the signature motif is demonstrated herein to apply across nuclear proteins as a class it is expected that different nuclear receptors display both coactivator and signature motif preferences that contribute to specificity of hormonal response (Ding *et al*, 1998, Molecular Endocrinology, 12, 302-313) which in turn points to selective pharmaceutical

intervention opportunities. Figures 1A and 5 below also indicate that individual motifs may differ in the strength to which they bind to nuclear receptors.

Note the NR box of Le Douarin *et al* (discussed above) did not disclose a signature motif within the meaning of the present invention because, for example, the NR box within the meaning of Le Douarin would be present in at most only 4 of the 39 signature motifs identified by the present invention in Figures 3A & 4 (see below). Furthermore, Le Douarin *et al* did not even suggest inhibitors of nuclear receptor - nuclear protein interaction.

The term "antibodies" is meant to include polyclonal antibodies, monoclonal antibodies, and the various types of antibody constructs such as for example $F(ab')_2$, Fab and single chain Fv. Antibodies are defined to be specifically binding if they bind with a K_a of greater than or equal to about 10^7 M^{-1}. Affinity of binding can be determined using conventional techniques, for example those described by Scatchard *et al.*, *Ann. N.Y. Acad. Sci.*, 51: 660 (1949).

Polyclonal antibodies can be readily generated from a variety of sources, for example, horses, cows, goats, sheep, dogs, chickens, rabbits, mice or rats, using procedures that are well-known in the art. In general, immunogen is administered to the host animal typically through parenteral injection. The immunogenicity may be enhanced through the use of an adjuvant, for example, Freund's complete or incomplete adjuvant. Following booster immunizations, small samples of serum are collected and tested for reactivity. Examples of various assays useful for such determination include those described in: *Antibodies: A Laboratory Manual*, Harlow and Lane (eds.), Cold Spring Harbor Laboratory Press, 1988; as well as procedures such as countercurrent immuno-electrophoresis (CIEP), radioimmunoassay, radioimmunoprecipitation, enzyme-linked immuno-sorbent assays (ELISA), dot blot assays, and sandwich assays, see U.S. Patent Nos. 4,376,110 and 4,486,530.

Monoclonal antibodies may be readily prepared using well-known procedures, see for example, the procedures described in U.S. Patent Nos. 4,902,614, 4,543,439 and 4,411,993; Monoclonal Antibodies, Hybridomas: *A New Dimension in Biological Analyses,* Plenum Press, Kennett, McKearn, and Bechtol (eds.), (1980).

Monoclonal antibodies can be produced using alternative techniques, such as those described by Alting-Mees *et al.*, "Monoclonal Antibody Expression Libraries: A Rapid Alternative to Hybridomas", *Strategies in Molecular Biology* 3: 1-9 (1990) which is

incorporated herein by reference. Similarly, binding partners can be constructed using recombinant DNA techniques to incorporate the variable regions of a gene that encodes a specific binding antibody. Such a technique is described in Larrick *et al.*, *Biotechnology*, 7: 394 (1989).

According to a further feature of the invention there is provided a pharmaceutical composition which comprises a novel inhibitor of the invention, or a pharmaceutically-acceptable salt thereof, in association with a pharmaceutically-acceptable diluent or carrier.

The composition may be in a form suitable for oral use, for example a tablet, capsule, aqueous or oily solution, suspension or emulsion; for topical use, for example a cream, ointment, gel or aqueous or oily solution or suspension; for nasal use, for example a snuff, nasal spray or nasal drops; for vaginal or rectal use, for example a suppository; for administration by inhalation, for example as a finely divided powder such as a dry powder, a microcrystalline form or a liquid aerosol; for sub-lingual or buccal use, for example a tablet or capsule; or for parenteral use (including intravenous, subcutaneous, intramuscular, intravascular or infusion), for example a sterile aqueous or oily solution or suspension. In general the above compositions may be prepared in a conventional manner using conventional excipients. For peptidic inhibitors, parenteral compositions are preferred.

The amount of active ingredient that is combined with one or more excipients to produce a single dosage form will necessarily vary depending upon the host treated and the particular route of administration. For example, a formulation intended for oral administration to humans will generally contain, for example, from 0.5 mg to 2 g of active agent compounded with an appropriate and convenient amount of excipients which may vary from about 5 to about 98 percent by weight of the total composition. Dosage unit forms will generally contain about 1 mg to about 500 mg of an active ingredient.

According to another aspect of the present invention there is provided a method of mapping nuclear receptor interaction domains in nuclear proteins in which the method comprises analysis of the sequence of a nuclear protein for the presence of signature motifs as defined herein in order to identify an interaction domain or a potential interaction domain. Preferably the analysis further comprises analysis of any potential interaction domains identified thereby for α-helicity and/or surface accessibility.

The invention is illustrated by the non-limiting Examples below in which. unless stated otherwise: temperatures are expressed in degrees Celsius; and peptide sequences are listed N-terminus to C-terminus.

Figures 1a/1b show the interaction of LXXLL motifs derived from coactivators with the ER.

Figure 1a: Yeast two hybrid interactions of LXXLL motifs, derived from the proteins RIP140, SRC1a and CBP with the LBDs of wild type or mutant ER. The sequences of the LXXLL motifs in the DNA binding domain (DBD) fusion proteins are indicated. DBD-LXXLL proteins were coexpressed with AAD-ER or AAD-ER Mut, which consist of an acidic activation domain (AAD) fused to the LBD of the wild-type ER, or a transcriptionally defective ER mutant, respectively. Reporter activities were determined in the presence or absence of 10^{-7}M 17-β-estradiol (E2) and expressed as units of β-galactosidase activity. Sequences listed in Figure 1a, from top to bottom, are listed as SEQ ID NO: 6-23 respectively.

Figure 1b: Effects of mutations in the RIP140 LXXLL motif located at amino acids 935-943 on binding of AAD-ER. Conserved leucine residues are boxed and mutated residues are circled. The reporter activity was determined in the presence (black bars) or absence (white bars) of 10^{-7}M E2. Sequences listed in Figure 1b, from top to bottom, are listed as SEQ ID NO: 24-32 respectively.

Figures 2a/2b/2c show that LXXLL motifs are required for binding of SRC1 to the ER LBD *in vitro* and for the ability of SRC1 to enhance ER activity *in vivo*.

Figure 2a: Wild type (SRC1a) and mutant (SRC1a-M1234) SRC1 proteins are shown schematically. The black bars represent the approximate locations of the LXXLL binding motifs in the linear SRC1a sequence and the shaded circles indicate the mutation of LXXLL binding motifs by replacement of conserved leucine residues with alanines (see Methods). Binding of wild type SRC1a or SRC1a-M1234 mutant to glutathione S transferase (GST) alone, to the ligand binding domain (aa 313-599) of ER (GST-AF2), or the SRC1 binding domain (aa 2058-2163) of CBP (GST-CBP) in the presence (+) and absence (-) of 10^{-6}M E2. The signals obtained with 10% of the input of [^{35}S] -labelled wild type and mutant SRC1 proteins are shown.

Figure 2b: The ability of increasing amounts of the peptides P-1 (SEQ ID NO: 2) and P-2 (SEQ ID NO: 72) to compete against the binding of wild type SRC1a to GST-AF2 in the presence of ligand is shown. The sequences of the P-1 and P-2 peptides are given at the foot of Fig 2b, and the conserved leucines and alanine substitutions are boxed.

Figure 2c: Wild type but not mutant SRC1e M123 potentiates activation by ER of the reporter gene 2ERE-pS2-CAT in transiently transfected Hela cells. Reporter activities obtained from extracts of transfected cells grown in the absence (white columns) or presence (black columns) of ligand (10^{-8}M E2). The amounts of ER, SRC1-wt and and SRC1-mut expression plasmids used in the transfections are indicated below the graph. The activities shown are averaged from duplicates.

Figures 3a/3b & 4 show that the LXXLL sequence is a signature motif in proteins that bind the LBDs of NRs.

Figure 3a: Alignment of LXXLL motif sequences present in human RIP140 [10], human SRC1a, mouse TIF2 [6], mouse CBP [23.24] , p300 [33], mouse TIF1 [11] and human TRIP proteins [12]. The conserved leucines are boxed and the amino acid numbers are given for each motif. Sequences listed in Figure 3A, from top to bottom, are listed as SEQ ID NO: 33-61 respectively.

Figure 3b: Schematic representation of the incidence of LXXLL motifs (black bars) in the sequences of proteins which bind NRs. The amino acid boundaries of the known NR binding sites are also shown.

Figure 4: Alignment of LXXLL motif sequences present in CBP, P300, p/CIP, ARA70 & TRIP 230. The conserved leucines are boxed and the amino acid numbers are given for each motif. Sequences listed in Figure 4, from top to bottom, are listed as SEQ ID NO: 62-71 respectively.

Figure 5 shows that the precise nature of the LXXLL signature motif affects the strength of the interaction between SRC-1a and the LBDs of ERα, ERβ and GR.

Figure 5a: The ERα LBD binds to the 4 signature motifs of SRC-1a with differing affinities in yeast 2-hybrid assays (SRC-1a motif 1-4 = SEQ ID 73-76). The order (decreasing affinity) is SRC-1a motif 2> SRC-1a motif 4 > SRC-1a motif 1> SRC-1a motif 3.

Figure 5b: The ERβ LBD binds to the signature motifs of SRC-1a with the same relative affinities seen with ERα. The order (decreasing affinity) is SRC-1a motif 2 > SRC-1a motif 4> SRC-1a motif 1 > SRC-1a motif 3.

Figure 5c: The GR LBD binds to the signature motifs of SRC-1 with differing affinities and the rank order of affinities differs to those seen with ERα and ERβ. The order (decreasing affinity) is SRC-a motif 4 > SRC-1a motif 1 = SRC-1a motif 2 = SRC-1a motif 3.

In Figure 5 the y-axis units represent relative β-galactosidase activity. In Figure 5 the x-axis motif numbering only indicates part of the actual sequence used which was as follows: 627-640, 684-696, 743-755 and 1428-1441.

The following abbreviations are used.

AAD	acidic activation domain
AF	activator function
DBD	DNA binding domain
E2	17-β-estradiol
ER	estrogen receptor
GR	glucocorticoid receptor
GST	glutathione S transferase
LBD	ligand binding domain
NR	nuclear receptor
PCR	polymerase chain reaction
RAR	retinoic acid receptor
RIP	receptor interacting protein
RXR	retinoid X receptor
SRC	steroid receptor coactivator
TIF	transcriptional intermediary factor
TRβ	thyroid hormone receptor β

Standard amino acid abbreviations have been used.

Alanine	Ala	A
Arginine	Arg	R
Asparagine	Asn	N
Aspartic Acid	Asp	D
Cysteine	Cys	C
Glutamic Acid	Glu	E
Glutamine	Gln	Q
Glycine	Gly	G
Histidine	His	H
Isoleucine	Ile	I
Leucine	Leu	L
Lysine	Lys	K
Methionine	Met	M
Phenylalanine	Phe	F
Proline	Pro	P
Serine	Ser	S
Threonine	Thr	T
Tryptophan	Trp	W
Tyrosine	Tyr	Y
Valine	Val	V
Any amino acid	Xaa	X

Point mutations will be referred to as follows: natural amino acid (using the 1 letter nomenclature) , position, new amino acid. For example "L636A" means that at position 636 a leucine (L) has been changed to alanine (A). Multiple mutations will be shown between square brackets.

Example 1

Mapping of interaction sites between Nuclear Receptor and Nuclear Protein

It has been previously demonstrated that the 140 kDa receptor interacting protein (RIP140) bound directly to NRs through at least two distinct sites located at the N- and C-termini of the protein [21]. To map these interaction sites in more detail, we examined a series of twenty different PCR-generated fragments of RIP140 coding sequence fused in frame with a heterologous DBD, for interaction with NRs in a two hybrid system. Remarkably, although the different constructs spanned the entire 1158 amino acids of RIP140 sequence, all but two displayed ligand-dependent interaction with ER, including five non-overlapping RIP140 sequences. By comparison of the sequences of the shortest interacting fragments we identified a short motif (LXXLL) common to all interacting fragments. In total, nine copies of the motif were identified in the RIP140 sequence, but the motif was absent in fragments showing no binding activity in our experiments.

To determine if these short sequences were sufficient to bind to NRs, we constructed a series of proteins consisting of a DBD fused to eight to ten amino acids incorporating one copy of each of the nine LXXLL motifs. As shown in **Fig.** 1a, each of the nine motifs present in RIP140 displayed strong ligand-dependent interaction with the LBD of the ER whereas the DBD alone showed no ability to bind. (Note that of the 10 motifs listed for RIP140, the 10th is a repeat of the 9th locus). Comparable results were obtained with the LBD of RAR (data not shown). Mutation of hydrophobic residues within H12 abolish AF2 activity and prevent the recruitment of RIP140 [10], TIF1 [11], TIF2 [6], SUG1 [13] and SRC1. Similarly, mutation of H12 residues M543 and L544 in the ER abolished the ligand-dependent interaction of all nine LXXLL motifs with ER (**Fig.** 1a). Taking these results together, we conclude that a short conserved motif comprised within as little as eight amino acids is sufficient to bind to transcriptionally active NRs. This discovery that such a relatively small motif can affect the interaction between two relatively large molecules is unprecedented in this field.

Secondary structure analysis using the Phd program [22] revealed that each of the nine copies of this motif in RIP140 occurred within a region predicted to be α-helical in nature, in which the conserved leucines would form a hydrophobic face.

Example 2

Mutational Analysis of a Signature Motif

To determine the sequence constraints required to observe a functional interaction, we carried out a partial mutational analysis of one of the RIP140 motifs (amino acids 935-943; **Fig.**1b). While western blot analysis showed no significant variation in the expression of the wild type and mutant fusion proteins (data not shown), mutation of valine 935 to alanine resulted in approximately ten fold reduction in the reporter activity in the presence of ligand which, when coupled with the observation that the first amino acid is hydrophobic in seven of the nine LXXLL motifs in RIP140, may indicate a preference for a hydrophobic residue at this position. Strikingly, mutation of any one of the three conserved leucine residues L936, L939 or L940 to alanine resulted in a complete loss of binding to the LBD of ER (**Fig.** 1b) and RAR (data not shown), emphasising their importance in mediating the interaction with NRs. In contrast, mutation to alanine of L941 (which is not conserved among the motifs; see **Fig.** 3a), had no effect on the ability of this sequence to bind to the ER LBD. Replacement of a conserved leucine residue with a valine was tolerated at L936, but not at L939 or L940 indicating that hydrophobic character alone is not sufficient to maintain an interaction with ER (Fig. 1b). The amino acids K937, Q938, S942 and E943 were not subjected to mutagenesis as they are not conserved among the motifs we have identified (see Fig. 3a).

Example 3

Analysis of Signature Motifs in Nuclear Proteins

The steroid receptor coactivator SRC1, which stimulates ligand-dependent transcriptional activity, was originally identified as a partial cDNA encoding a protein capable of interacting with the progesterone receptor by means of a 196 amino acid C-terminal region [3]. We noted that the eight most C-terminal amino acids fit the LXXLL consensus, and indeed this sequence (DBD-SRC1a 1434-1441) displayed strong ligand-induced binding to ER, but not the ER H12 mutant (Fig. 1a). Subsequent studies have identified full-length SRC1 (SRC1a) from mouse (1459 amino acids) [4,5] and human (1441 amino acids) tissues. Both murine and human SRC1a proteins interact with multiple NRs, and contain an additional interaction region between residues 569 to 789 [5] and 570-780, respectively. Three copies of the LXXLL motif were identified in this central interaction domain of human SRC1a (see Fig.

3a & 3b), each of which displayed ligand-dependent binding to both ER (Fig. 1a) and RAR (data not shown) in the two hybrid assay, but not the ER H12 mutant. Interestingly, the sequences and relative positions of the three motifs in the central domain of SRC1a are conserved in the related coactivator protein Transcriptional Intermediary Factor 2 (TIF2) (Fig. 3a + b), and correspond to the region of TIF2 known to bind to NRs [6]. However, unlike SRC1a, TIF2 appears to lack a motif at its C-terminus. In addition, we noted that SRC1 contains three other sequences matching the LXXLL consensus. Although the motif at residues 45-53 is predicted by the Phd program to be α-helical and lies within the basic helix-loop-helix domain at the N-terminus of SRC1a [5], it showed only a very weak (6-fold) interaction with liganded ER (Fig. 1a) or RAR (not shown) in the yeast two hybrid assay. This is consistent with the observed absence of strong NR-binding activity associated with the N-terminus of SRC1. The other two motifs within residues 111-118 and 912-920 both contain proline residues and are unlikely to adopt α-helical structure according to the Phd program. Indeed, these sequences showed no detectable interaction with NRs in our binding assays (Fig. 1a), which strongly suggests a preference for appropriate secondary structure for binding of LXXLL sequences to NRs.

Recent reports have indicated that CBP/p300 proteins, which were originally identified as coactivators for CREB [23,24], are coactivators for many transcription factors including NRs [4,5,8,9] and may serve as integrators of several signalling pathways [4]. CBP was shown to bind directly to NRs via its N-terminal 101 amino acids [4], with a possible RXR-specific binding site between residues 356-495 [8]. Our analysis showed that the CBP sequence harbours copies of the LXXLL motif within positions 68-78, and 356-364, which are conserved in the p300 sequence (amino acids 80-90 and 341-351; Fig 3a). Indeed, when tested in the two hybrid assay, the N-terminus of CBP (amino acids 1-101; data shown) and the LXXLL motif at residues 68-75 of the CBP sequence (Fig. 1a) displayed ligand-dependent binding to ER, but not the transciptionally defective ER mutant (Fig. 1a).

Example 4

The Binding Of Coactivator Proteins To NRs Is Dependent On Signature Motifs

To demonstrate that the binding of coactivator proteins to NRs is dependent on LXXLL motifs, we introduced alanine substitutions in SRC1a at the conserved leucine

couplets at residues [L636A, L637A, L693A, L694A, L752A, L753A, L1438A, L1439A] thus effectively creating a mutant protein (SRC1a-M1234) in which all the four functional binding motifs were disabled. We then compared the ability of *in vitro* translated SRC1a and SRC1a-M1234 to bind to the ligand binding domain of the mouse estrogen receptor fused to glutathione-S-transferase (GST-AF2) in GST pulldown experiments (Maniatis *et al.*,(1982) Molecular Cloning. A Laboratory Manual. Cold Spring Harbour Laboratory, Cold Spring Harbour, New York.). As shown in Fig. 2a, while wild type SRC1a protein displayed ligand-dependent binding to GST-AF2, SRC1a-M1234 failed to bind to GST-AF2 either in the presence or absence of ligand. To confirm that the mutations did not induce gross structural disruption of the SRC1a-M1234, we compared the ability of the *in vitro* translated proteins to interact with amino acids 2058-2163 of CBP, which was previously defined as the SRC1 binding domain [4]. Both proteins retained strong binding to GST-CBP (Fig. 2a) indicating that this SRC1 function remained intact in both wild type and mutant proteins. In addition we showed that the binding of wild type SRC1a to GST-AF2 was competed by increasing concentrations of a short peptide (P1) corresponding to the motif at the C-terminus of SRC1a (Fig. 2b). In contrast, a similar peptide (P2) in which the LXXLL motif was mutated, or peptides unrelated to the LXXLL motif (data not shown), did not compete the binding of SRC1a to GST-AF2 (Fig. 2b).

Finally, to demonstrate that LXXLL motifs are necessary for the function of SRC1 *in vivo,* we compared the abilities of wild type SRC1 and a mutant protein in which all LXXLL motifs were disabled to enhance the activity of mouse ER in transient transfection experiments. As shown in Fig. 2c, wild type SRC1 enhanced the activity of ER in a concentration-dependent manner. In contrast, the SRC1 mutant, which was unable to bind ER (Fig. 2a) had no stimulatory effect, but reduced ER activity by up to 50% at the highest concentration (Fig. 2c). This apparent dominant negative property of the mutant SRC1 is likely due to its ability to maintain interactions with CBP while failing to interact with NRs (Fig. 2a). This result is of interest given the recent evidence that SRC1 and CBP/p300 may exist as a complex *in vivo* [9], and that CBP also has NR binding activity [4, 8], as our data suggest that the interactions between NRs and CBP are insufficient to compensate for the inability of the SRC1a mutant protein to bind NRs, at least under these conditions. It remains

to be determined whether NRs are engaged simultaneously by p160 and p300 proteins functioning independently or as a complex.

Examination of the sequences of other proteins known to bind to NRs revealed them to contain one or more copies of the LXXLL motif. TIF1 contains a single motif (residues 722-732) within the minimal region known to be required for its interaction with NRs [11, 25]. The truncated proteins TRIPs2-5, TRIP8 and TRIP9, which were isolated in a two hybrid screen for TR-interacting proteins [12], each contain at least one copy of the LXXLL motif (Fig. 3a), whereas the motif was absent in TRIPs whose interaction with TR was ligand-independent. An alignment of a selection of these sequences is shown in Fig. 3a, while Fig. 3b shows the incidence of motifs in the sequences of RIP140, SRC1a, TIF1, TIF2, CBP and p300, and the boundaries of known receptor interaction domains in these proteins. Interestingly, motifs were also identified in several other proteins for which evidence exists of interaction with NRs, including Ara70 [26], SW13 [27], and the RelA (p65) subunit of NFκ-B [28], although the receptor interaction domains in these proteins have not been mapped. The ability of other proteins containing LXXLL motifs to bind to NRs will depend on their subcellular localisation, as well as the α-helicity and surface accessibility of the motifs. While it is clear that the conserved leucine residues are essential for the function of the motif, other amino acids may also be important given the degree of sequence conservation of equivalent motifs in SRC1/TIF2 or CBP/p300.

As many NR binding proteins contain multiple copies of the LXXLL motif it remains to be established whether this facilitates the simultaneous contact of individual partners in homo- and heterodimers of NRs, or whether it serves to provide alternative interaction surfaces to accomodate conformational changes imposed by the binding of NRs to different response elements. The systematic mutation of LXXLL motifs in coactivators such as SRC1 and CBP may allow us to decouple crosstalk or synergy between different signal transduction pathways, and thus provide a better understanding of their proposed roles as coactivators and integrators.

Example 5

Two Hybrid Interaction Assays

The yeast reporter strain used for all two hybrid assays was W303-1B (HMLα MATα HMRa his3-11, 15 trp1-1 ade2-1 can1-100 leu2-3, 11, ura3) carrying the plasmid pRLΔ21-U3ERE which contains a lacZ reporter gene driven by three estrogen response elements (EREs) [29]. The plasmids pBL1 and pASV3 which express the human ER DNA binding domain (DBD) and the VP16 acidic activation domain (AAD) respectively [30], were used to generate DBD or AAD fusion proteins for two hybrid interaction analyses. DBD-LXXLL motif fusion proteins were generated by ligation of phosphorylated, annealed oligonucleotide pairs into the pBL1 vector. AAD-ER was constructed by cloning a PCR fragment encoding amino acids 282-595 of the human ER into pASV3. AAD-ER Mut was constructed in a similar fashion except that the amino acids M543 and L544 or ER were mutated to alanines by recombinant PCR. All fusion constructs were fully sequenced. Transformants containing the desired plasmids were obtained by selection for the appropriate plasmid markers and were grown to late log phase in 15 ml of selective medium (yeast nitrogen base containing 1% glucose and appropriate supplements) in the presence or absence of 10^{-7}M 17-β-estradiol (E2).

The expression of DBD- and AAD- fusion proteins in yeast cell-free extracts was verified by immunodetection using a monoclonal antibody recognising the human ER (a gift from P. Chambon, Strasbourg). The antibody recognises the "F" region of the LBD in the human ER, and also the "F" region tag at the N-termini of the DBD fusion proteins [30]. Equal amounts of protein were electrophoresed on polyacrylamide gels and transferred to nitrocellulose for western blotting. The preparation of cell-free extracts by the glass bead method and the measurement of β-galactosidase activity in the extracts were performed as previously described [29]. Two hybrid experiments were repeated several times, and the data shown in Figs. 1a and 1b represent reporter activities as measured in a single representative experiment. The β-galactosidase activities are expressed as nmoles/minute/μg protein.

Example 6

***In Vitro* Binding And Peptide Inhibition Assays**

GST-AF2 consists of the ligand binding domain of the mouse ER (amino acids 313-599) fused to glutathione-S-transferase and has been described previously [31]. GST-CBP consists of GST fused to the SRC1-binding domain of CBP and was constructed by cloning a PCR fragment encoding residues 2058-2163 of mouse CBP into the vector pGEX2TK (Pharmacia). Human SRC1a and SRC1e cDNAs were isolated form a human B cell cDNA library and cloned into a modified version of the expression vector pSG5. SRC1a M1234 and SRC1eM123 were constructed by recombinant PCR to introduce the mutations [L636A, L637A, L693A, L694A, L752A, L753A, L1438A, L1439A] or [L636A, L637A, L693A, L694A, L752A, L753A] respectively. All SRC1 constructs were fully sequenced. GST-SEPHAROSE™ beads were loaded with GST alone or GST-fusion proteins prepared from bacterial cell-free extracts. [^{35}S]-labelled SRC1 proteins were generated by *in vitro* translation and tested for interaction with GST proteins in the presence or absence of 10^{-6}M estradiol (E2) as previously described [21]. Binding was carried out for 3 hours at 4° with gentle mixing in NETN buffer (100 mM NaCl, 1 mM EDTA, 0.5 % NP-40, 20 mM Tris HCl, pH 8.0) containing protease inhibitors in a final volume of 1 ml. Peptides P-1 and P-2 were dissolved in water at a concentration of 4mg/ml and added individually to GST-binding reactions immediately before the addition of ligand. The increasing amounts of peptide added in the competition experiments shown corresponded to 2.5, 5, 12.5 and 25 μM.

Using analogous methodology, peptides KLVQLLTTT (SEQ ID NO: 3), ILHRLLQE (SEQ ID NO: 4) and LLQQLLTE (SEQ ID NO: 5) can be shown to be inhibitors.

Example 7

Transient Reporter Assays

Hela cells were transfected with 1μg of reporter 2ERE-pS2-CAT [32], 150 ng of β-galactosidase expression plasmid (internal control), 10 ng of ER expression plasmid and 50 or 200 ng of SRC1 expression plasmids or empty vector per well (in duplicate) using 24-well plates. Transfected cells were incubated overnight in Dulbecco's modified Eagle's medium without phenol red and containing 10% charcoal-treated FBS, and washed in fresh medium before addition of ligand (10^{-8}M E2) or vehicle. After 40 hrs, cells were harvested and

extracts analysed for CAT and β-galactosidase activities [14,21]. β-galactosidase activities were used to correct for differences in transfection efficiency.

Example 8

Pharmaceutical Composition

The following illustrates a representative pharmaceutical dosage form containing a peptide inhibitor and which may be used for therapy.

Injectable solution

A sterile aqueous solution, for injection, containing per ml of solution:

Peptide P-1	5.0mg
Sodium acetate trihydrate	6.8mg
Sodium chloride	7.2mg
Tween 20	0.05mg

A typical dose of peptide for adult humans is 30mg.

Example 9

The strength of the interaction of coactivator signature motifs to NRs varies depending on the precise motif sequence

The interaction between the ERα LBD and a range of different LXXLL motif fusion proteins appeared to yield different reporter gene activity indicating that the ERα LBD interacted with the LXXLL motifs with differing affinities (see Example 5), therefore the strength of these interactions was investigated more closely. The motifs of SRC-1a were tested for their relative interaction specificities with the glucocorticoid receptor and estrogen receptor isoforms α and β in a yeast two hybrid assay.

The SRC-1a motifs 1-4 (SEQ ID 73-76) were expressed as fusion proteins with the LexA DNA binding domain. Motif fusion proteins were generated by ligation of annealed oligonucleotide pairs in frame into the ADH promoter driven LexA DBD vector YCp14-ADH-LexA. YCp14ADH1-LexA is a plasmid from which LexA is expressed under control of

the *S. cerevisiae* ADH1 promoter. The backbone of this vector is the plasmid RS314 (Sikorski and Hieter, 1989). Between the Sac I and Kpn I restriction enzyme sites in the polylinker of this vector we have placed an expression cassette comprising the promoter of the *S. cerevisiae* ADH1 gene, the coding region of the *E. coli* LexA gene and the transcription termination region region of the *S. cerevisiae* ADH1 gene. The promoter region of ADH1 consists of a 1.4kb Bam HI- Hind III fragment from the plasmid pADNS (Colicelli *et al*, 1989). Following the Hind III site is the coding region (amino acids 1-202) of *E. coli* LexA (Horii *et al*, 1981; Miki *et al*, 1981; Markham *et al*, 1981). The *E. coli* LexA fragment was obtained as a Hind III- Pst I fragment from the plasmid pBXL1 (Martin *et al*, 1990). This sequence corresponds to nucleotides 95-710 of Genbank entry g146607. Following this sequence is a region which encodes a polylinker (SEQ ID 77).

GAATTCCTGCAGCCCGGGGTCGACACTAGTTAACTAGCGGCCGC

This polylinker adds the amino acids EFLQPGVDTS (SEQ ID NO: 80) to the carboxy terminus of LexA. The Not I site at the end of this linker is linked to a DNA fragment which includes the transcription terminator region of the *S. cerevisiae* ADH1 gene. This fragment is the 0.6kb Not I-Bam HI from the plasmid pADNS (Colicelli *et al*, 1989).

The NR LBDs were expressed as fusions with the Gal4 transcriptional AD. The LBD fusions were constructed by cloning a PCR fragment encoding the amino acids corresponding to the LBDs into YCp15Gal1-r11.YCp15Gal1-r11 is a plasmid from which fusions of the activation region of the *S. cerevisiae* Gal4 protein may be expressed under the control of the *S. cerevisiae GAL1* promoter. The backbone of this vector is the plasmid RS315 (Sikorski and Hieter, 1989). Between the Sac I and Kpn I restriction enzyme sites in the polylinker of this vector we have placed an expression cassette comprising the promoter of the *S. cerevisiae GAL1* gene, the coding region of the fusion protein and the transcriptional termination region of the *S. cerevisiae* ADH1 gene. The *GAL1* promoter (Johnson and Davis, 1984;Yocum *et al*, 1984; West *et al*, 1984) was obtained by amplification of a *S. cerevisiae* genomic fragment by the polymerase chain reaction (PCR). This fragment corresponds to nucleotides 177 to 809 of Genbank database entry g171546. This is followed by the sequence

AAGCTTCCACCATGGTGCCAAAGAAGAAACGTAAAGTT (SEQ ID 78).

This sequence provides a translation initiation codon and a sequence which encodes the amino acids MVPKKKRKV (SEQ ID NO: 81). The last seven residues of this peptide

correspond to a region identified as a nuclear localisation signal in the SV40 T antigen (amino acids 126-132 of Genbank entry g310678; Fiers *et al*, 1978; Reddy *et al*, 1978). This region is linked to sequence encoding the region II transcription activation domain (amino acids 768-881) of Gal4, as defined by Ma and Ptashne (1987). The sequence was isolated by PCR from plasmid pBXGalII (P. Broad unpublished) which is a mammalian version of the yeast expression vector pMA236 (Ma and Ptashne, 1987) and corresponds to nucleotides 2744 to 3085 of Genbank entry g171557 (Laughon and Gesteland (1984). This sequence is followed by the polylinker

TCTAGACTGCAGACTAGTAGATCTCCCGGGGCGGCCGC (SEQ ID 79).

All fusion constructs were fully sequenced. The vectors YCp14-ADH-lexA and YCp15Gal1-r11 replicate as a single copy plasmids in yeast(ARS-CEN) and have TRP1 and LEU2 markers respectively.

The *S. cerevisiae* strain MEY132 (M. Egerton, Zeneca Pharmaceuticals, unpublished, genotype (*Mata leu2-3,112 ura3-52 trp1 his4 rme1*) was employed as a host strain. A reporter gene consisting of the E. coli lacZ (β-galactosidase) gene under the control of a promoter containing two binding sites for the lexA protein was integrated at the *ura3* locus of this strain. The reporter gene plasmid, JP159-lexRE was constructed using plasmid JP159 as a backbone (J. Pearlberg, Ph.D. Thesis (1994) Harvard University). This is a shuttle vector which contains the *S. cerevisiae URA3* as a marker. The plasmid contains the the E.coli β-galactosidase gene under the control of a minimal promoter containing the TATA box and transcription initiation site from the *S. cerevisiae GAL1* promoter (Dixon *et al* 1997). Upstream of this promoter is a terminator from the *GAL11* gene. Between the Xba I and Sal I sites in the promoter of this plasmid a 35 nucleotide sequence corresponding to a naturally occurring binding sites for the LexA protein in the promoter of the colicin E1 gene (Ebina *et al*, 1983) the sequence corresponds to residues 20-54 of Genbank entry g144345). This sequence contains two LexA operators and is therefore referred to as "2lex". This reporter plasmid was linearised within the *URA3* gene and integrated into the *ura3* locus of MEY132 to give the yeast strain MEY132-lexRE

Transformants containing the 2 hybrid fusion constructs were grown to late log phase in 20ml selective medium (2% glucose and appropriate supplements). They were then diluted into 2% galactose containing medium, in the presence or absence of ligand. As controls each

LBD fusion was coexpressed with the LexA DBD lacking the coactivator motifs and similarly each motif LexA DBD fusion was coexpressed with Gal4 AD lacking the NR LBD. The relative expression levels of DBD fusion proteins were determined by immunodetection using a monoclonal antibody which recognises LexA.

The estrogen receptor isoforms α and β relative interaction specificities for the 4 motifs of SRC-1a are ranked as follows, 2>4>1>3 (see Figure 5) . Glucocorticoid receptor specificities are as follows 4>1=2=3.

The yeast 2-hybrid system used in this Example utilises a single copy integrated reporter gene construct. This enables a quantitative comparison between the different yeast strains bearing the different SRC1a motifs. The data presented in Figure 5 suggest that motif 3 (SRC 1a, 748-753) interacts very weakly, using this system, with ER. Longer exposures to oestradiol (not shown) do however reveal a significant interaction. It is noted that Figure 1A herein clearly shows an interaction between motif 3 and ER but this interaction is significantly weaker than that seen with the other motifs. Any differences in this regard between Figures 5 and 1A can be explained by experimental design features. For example, vectors used to generate the Figure 5 data were low copy number (centromere containing) vectors, whereas vectors used to generate the Figure 1A data were multicopy (2 micron) vectors.

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SEQUENCE LISTING

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(ii) TITLE OF INVENTION: Chemical Compounds

(iii) NUMBER OF SEQUENCES: 79

(iv) COMPUTER READABLE FORM:
(A) MEDIUM TYPE: Floppy disk
(B) COMPUTER: IBM PC compatible
(C) OPERATING SYSTEM: PC-DOS/MS-DOS
(D) SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)

(vi) PRIOR APPLICATION DATA:
(A) APPLICATION NUMBER: GB 9708676.3
(B) FILING DATE: 30-APR-1997

(2) INFORMATION FOR SEQ ID NO: 1:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 5 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:

Leu Xaa Xaa Leu Leu
1 5

(2) INFORMATION FOR SEQ ID NO: 2:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 14 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:

Pro Gln Ala Gln Gln Lys Ser Leu Leu Gln Gln Leu Leu Thr
1 5 10

(2) INFORMATION FOR SEQ ID NO: 3:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3:

Lys Leu Val Gln Leu Leu Thr Thr Thr
1 5

(2) INFORMATION FOR SEQ ID NO: 4:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 4:

Ile Leu His Arg Leu Leu Gln Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 5:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:

Leu Leu Gln Gln Leu Leu Thr Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 6:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 6:

Tyr Leu Glu Gly Leu Leu Met His Gln Ala
1 5 10

(2) INFORMATION FOR SEQ ID NO: 7:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 7:

Leu Leu Ala Ser Leu Leu Gln Ser
1 5

(2) INFORMATION FOR SEQ ID NO: 8:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 8:

His Leu Lys Thr Leu Leu Lys Lys Ser
1 5

(2) INFORMATION FOR SEQ ID NO: 9:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9:

Gln Leu Ala Leu Leu Leu Ser Ser
1 5

(2) INFORMATION FOR SEQ ID NO: 10:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 10:

Leu Leu Leu His Leu Leu Lys Ser Gln
1 5

(2) INFORMATION FOR SEQ ID NO: 11:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 11:

Val Thr Leu Leu Gln Leu Leu Leu Gly
1 5

(2) INFORMATION FOR SEQ ID NO: 12:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:

Val Leu Gln Leu Leu Leu Gly Asn
1 5

(2) INFORMATION FOR SEQ ID NO: 13:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 13:

Leu Leu Ser Arg Leu Leu Arg Gln
1 5

(2) INFORMATION FOR SEQ ID NO: 14:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 14:

Val Leu Lys Gln Leu Leu Leu Ser Glu Asn
1 5 10

(2) INFORMATION FOR SEQ ID NO: 15:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 15:

Val Leu Lys Gln Leu Leu Leu Ser
1 5

(2) INFORMATION FOR SEQ ID NO: 16:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 16:

Glu Leu Ala Glu Leu Leu Ser Ala Asn
1 5

(2) INFORMATION FOR SEQ ID NO: 17:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 17:

Ser Leu Gly Pro Leu Leu Leu Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 18:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 18:

Lys Leu Val Gln Leu Leu Thr Thr Thr
1 5

(2) INFORMATION FOR SEQ ID NO: 19:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 19:

Ile Leu His Arg Leu Leu Gln Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 20:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 20:

Leu Leu Arg Tyr Leu Leu Asp Lys
1 5

(2) INFORMATION FOR SEQ ID NO: 21:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 21:

Gln Leu Asp Glu Leu Leu Cys Pro Pro
1 5

(2) INFORMATION FOR SEQ ID NO: 22:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 22:

Leu Leu Gln Gln Leu Leu Thr Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 23:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 23:

Gln Leu Ser Glu Leu Leu Arg Gly
1 5

(2) INFORMATION FOR SEQ ID NO: 24:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 24:

Val Leu Lys Gln Leu Leu Leu Ser Glu Asn
1 5 10

(2) INFORMATION FOR SEQ ID NO: 25:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 25:

Ala Leu Lys Gln Leu Leu Leu Ser Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 26:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 26:

Val Ala Lys Gln Leu Leu Leu Ser Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 27:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 27:

Val Val Lys Gln Leu Leu Leu Ser Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 28:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 28:

Val Leu Lys Gln Ala Leu Leu Ser Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 29:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 29:

Val Leu Lys Gln Val Leu Leu Ser Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 30:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 30:

Val Leu Lys Gln Leu Ala Leu Ser Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 31:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 31:

Val Leu Lys Gln Leu Val Leu Ser Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 32:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 32:

Val Leu Lys Gln Leu Leu Ala Ser Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 33:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 33:

Tyr Leu Glu Gly Leu Leu Met His Gln Ala Ala
1 5 10

(2) INFORMATION FOR SEQ ID NO: 34:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 34:

Leu Leu Ala Ser Leu Leu Gln Ser Glu Ser Ser
1 5 10

(2) INFORMATION FOR SEQ ID NO: 35:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35:

His Leu Lys Thr Leu Leu Lys Lys Ser Lys Val
1 5 10

(2) INFORMATION FOR SEQ ID NO: 36:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 36:

Gln Leu Ala Leu Leu Leu Ser Ser Glu Ala His
1 5 10

(2) INFORMATION FOR SEQ ID NO: 37:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 37:

Leu Leu Leu His Leu Leu Lys Ser Gln Thr Ile
1 5 10

(2) INFORMATION FOR SEQ ID NO: 38:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 38:

Leu Leu Gln Leu Leu Leu Gly His Lys Asn Glu
1 5 10

(2) INFORMATION FOR SEQ ID NO: 39:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 39:

Val Leu Gln Leu Leu Leu Gly Asn Pro Lys Gly
1 5 10

(2) INFORMATION FOR SEQ ID NO: 40:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 40:

Leu Leu Ser Arg Leu Leu Arg Gln Asn Gln Asp
1 5 10

(2) INFORMATION FOR SEQ ID NO: 41:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 41:

Val Leu Lys Gln Leu Leu Leu Ser Glu Asn Cys
1 5 10

(2) INFORMATION FOR SEQ ID NO: 42:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42:

Lys Leu Val Gln Leu Leu Thr Thr Thr Ala Glu
1 5 10

(2) INFORMATION FOR SEQ ID NO: 43:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 43:

Ile Leu His Arg Leu Leu Gln Glu Gly Ser Pro
1 5 10

(2) INFORMATION FOR SEQ ID NO: 44:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 44:

Leu Leu Arg Tyr Leu Leu Asp Lys Asp Glu Lys
1 5 10

(2) INFORMATION FOR SEQ ID NO: 45:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 45:

Leu Leu Gln Gln Leu Leu Thr Glu
1 5

(2) INFORMATION FOR SEQ ID NO: 46:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 46:

Lys Leu Leu Gln Leu Leu Thr Thr Lys Ser Asp
1 5 10

(2) INFORMATION FOR SEQ ID NO: 47:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 47:

Ile Leu His Arg Leu Leu Gln Asp Ser Ser Ser
1 5 10

(2) INFORMATION FOR SEQ ID NO: 48:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 48:

Leu Leu Arg Tyr Leu Leu Asp Lys Asp Asp Thr
1 5 10

(2) INFORMATION FOR SEQ ID NO: 49:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 49:

Gln Leu Ser Glu Leu Leu Arg Gly Gly Ser Gly
1 5 10

(2) INFORMATION FOR SEQ ID NO: 50:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 50:

Gln Leu Val Leu Leu Leu His Ala His Lys Cys
1 5 10

(2) INFORMATION FOR SEQ ID NO: 51:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 51:

Gln Leu Ser Glu Leu Leu Arg Gly Ser Ser Pro
1 5 10

(2) INFORMATION FOR SEQ ID NO: 52:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 52:

Gln Leu Val Leu Leu Leu His Ala His Lys Cys
1 5 10

(2) INFORMATION FOR SEQ ID NO: 53:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 53:

Ile Leu Thr Ser Leu Leu Leu Asn Ser Ser Gln
1 5 10

(2) INFORMATION FOR SEQ ID NO: 54:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 54:

Met Leu Met Asn Leu Leu Lys Asp Asn Pro Ala
1 5 10

(2) INFORMATION FOR SEQ ID NO: 55:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 55:

Thr Leu Arg Ser Leu Leu Leu Asn Pro His Leu
1 5 10

(2) INFORMATION FOR SEQ ID NO: 56:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 56:

Arg Leu Ala Val Leu Leu Pro Gly Arg His Pro
1 5 10

(2) INFORMATION FOR SEQ ID NO: 57:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 57:

Glu Leu His Asn Leu Leu Glu Val Val Ser Gln
1 5 10

(2) INFORMATION FOR SEQ ID NO: 58:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 58:

Thr Leu Arg Asp Leu Leu Thr Thr Thr Ala Gly
1 5 10

(2) INFORMATION FOR SEQ ID NO: 59:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 59:

Phe Leu Asp Phe Leu Leu Gly Phe Ser Ala Gly
1 5 10

(2) INFORMATION FOR SEQ ID NO: 60:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 60:

Val Leu Glu Leu Leu Leu Arg Ala Gly Ala Asn
1 5 10

(2) INFORMATION FOR SEQ ID NO: 61:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 61:

Ile Leu Ala Arg Leu Leu Arg Ala His Gly Ala
1 5 10

(2) INFORMATION FOR SEQ ID NO: 62:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:

Ala Leu Gln Asp Leu Leu Arg Thr Leu Lys Ser
1 5 10

(2) INFORMATION FOR SEQ ID NO: 63:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 63:

Ala Leu Gln Asn Leu Leu Arg Thr Leu Arg Ser
1 5 10

(2) INFORMATION FOR SEQ ID NO: 64:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 64:

Lys Leu Leu Gln Leu Leu Thr Cys Ser Ser Asp
1 5 10

(2) INFORMATION FOR SEQ ID NO: 65:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 65:

Ile Leu His Lys Leu Leu Gln Asn Gly Asn Ser
1 5 10

(2) INFORMATION FOR SEQ ID NO: 66:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 66:

Leu Leu Arg Tyr Leu Leu Asp Arg Asp Asp Pro
1 5 10

(2) INFORMATION FOR SEQ ID NO: 67:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 67:

Gln Leu Tyr Ser Leu Leu Gly Gln Phe Asn Cys
1 5 10

(2) INFORMATION FOR SEQ ID NO: 68:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 68:

Glu Leu Glu Asn Leu Leu Gln Gln Gly Gly
1 5 10

(2) INFORMATION FOR SEQ ID NO: 69:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 69:

Val Leu Gln Lys Leu Leu Lys Glu Lys Asp
1 5 10

(2) INFORMATION FOR SEQ ID NO: 70:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 70:

Glu Leu Asn Gln Leu Leu Asn Ala Val Lys
1 5 10

(2) INFORMATION FOR SEQ ID NO: 71:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 71:

Val Leu Lys Asp Leu Leu Lys Gln
1 5

(2) INFORMATION FOR SEQ ID NO: 72:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 14 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 72:

Pro Gln Ala Gln Gln Lys Ser Leu Leu Gln Gln Ala Ala Thr
1 5 10

(2) INFORMATION FOR SEQ ID NO: 73:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 14 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 73:

Ser Gln Thr Ser His Lys Leu Val Gln Leu Leu Thr Thr Thr
1 5 10

(2) INFORMATION FOR SEQ ID NO: 74:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 13 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 74:

Thr Ala Arg His Lys Ile Leu His Arg Leu Leu Gln Glu
1 5 10

(2) INFORMATION FOR SEQ ID NO: 75:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 13 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 75:

Ser Lys Asp His Gln Leu Leu Arg Tyr Leu Leu Asp Lys
1 5 10

(2) INFORMATION FOR SEQ ID NO: 76:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 14 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 76:

Gln Ala Gln Gln Lys Ser Leu Leu Gln Gln Leu Leu Thr Glu
1 5 10

(2) INFORMATION FOR SEQ ID NO: 77:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 44 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: other nucleic acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 77:

GAATTCCTGC AGCCCGGGGT CGACACTAGT TAACTAGCGG CCGC 44

(2) INFORMATION FOR SEQ ID NO: 78:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 38 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: other nucleic acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 78:

AAGCTTCCAC CATGGTGCCA AAGAAGAAAC GTAAAGTT 38

(2) INFORMATION FOR SEQ ID NO: 79:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 38 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: other nucleic acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 79:

TCTAGACTGC AGACTAGTAG ATCTCCCGGG GCGGCCGC 38

(2) INFORMATION FOR SEQ ID NO: 80:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 80:

Glu Phe Leu Gln Pro Gly Val Asp Thr Ser
1 5 10

(2) INFORMATION FOR SEQ ID NO: 81:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9 amino acids
(B) TYPE: amino acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: peptide

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 81:

Met Val Pro Lys Lys Lys Arg Lys Val
1 5

CLAIMS

1. A method for identifying inhibitor compounds capable of reducing the interaction between:

a) a first region which is a signature motif on a nuclear protein, and

b) a second region which is that part of a nuclear receptor which is capable of interacting with the nuclear protein through binding to the signature motif,

wherein:

the nuclear protein is a bridging factor that is responsible for the interaction between a liganded nuclear receptor and a transcription initiation complex involved in regulation of gene expression;

the nuclear receptor is a transcription factor;

the signature motif is a short sequence of amino acid residues which is the key structural element of a nuclear protein which binds to a liganded nuclear receptor as part of the process of the activation or repression of target genes; and

in which the method comprises taking:

i) the potential inhibitor compound;

ii) the liganded nuclear receptor or a fragment thereof in which the fragment comprises the second region defined in this claim in b) above;

iii) a fragment comprising a signature motif of the nuclear protein; and

iv) detecting the presence or absence of inhibition of the interaction between ii) and iii).

2 A method according to claim 1 in which the signature motif is B^1XXLL in which B^1 is any natural hydrophobic amino acid, L is leucine and X independently represents any natural amino acid.

3 A method according to claim 2 in which B^1 is leucine or valine.

4 A method according to claim 3 in which B^1 is leucine.

5 A method according to any one of claims 2-4 in which the signature motif is further defined as B^2B^1XXLL wherein B^2 is a hydrophobic amino acid.

6 A method according to claim 5 in which B^2 is selected from the group consisting of isoleucine, leucine, methionine, phenylalanine, tryptophan, tyrosine and valine.

7. A method according to any one of claims 1-6 in which the nuclear protein is a coactivator.

8. A method according to claim 7 in which the coactivator is selected from the group consisting of RIP 140, SRC-1, TIF2, CBP, p300, TIF1, Trip1, Trip2, Trip3, Trip4, Trip5, Trip8, Trip9, p/CIP, ARA70 & Trip230.

9. A method according to any one of claims 1-6 in which the transcription factor is a steroid hormone receptor.

10. A method according to claim 9 in which the steroid hormone receptor is selected from the group consisting of oestrogen receptor, progesterone receptor, androgen receptor and glucocorticoid receptor.

11. A method according to claim 10 in which the steroid hormone receptor is oestrogen receptor.

12. A method according to any preceding claim wherein the method is in the form of a 2-hybrid assay system.

13 A method according to any preceding claim wherein the potential inhibitor is in the form of a peptide library based on a signature motif as defined in any one of claims 2-6.

14. A novel inhibitor identified according to the method defined in any one of claims 1-13 which reduces the interaction between

a) a first region which is a signature motif on a nuclear protein, and

b) a second region which is that part of a nuclear receptor which is capable of interacting with the nuclear protein through binding to the signature motif,

wherein:

the nuclear protein is a bridging factor that is responsible for the interaction between a liganded nuclear receptor and the transcription initiation complex involved in regulation of gene expression;

the nuclear receptor is a transcription factor;

the signature motif is a short sequence of amino acid residues which is the key structural element of a nuclear protein which binds to a liganded nuclear receptor as part of the process of the activation or repression of target genes.

15 An inhibitor according to claim 14 which is a peptide of less than 15 amino acid residues comprising the signature motif defined in any one of claims 1-6.

16 An inhibitor according to claim 15 selected from the group consisting of PQAQQKSLLQQLLT (SEQ ID NO: 2), KLVQLLTTT (SEQ ID NO: 3), ILHRLLQE (SEQ ID NO: 4) and LLQQLLTE (SEQ ID NO: 5).

17 An inhibitor according to claim 14 comprising an antibody which specifically binds to a signature motif on a nuclear protein.

18 A pharmaceutical composition which comprises an inhibitor as defined in any one of claims 14-17 or a pharmaceutically-acceptable salt thereof, in association with a pharmaceutically-acceptable diluent or carrier.

19 A method of mapping nuclear receptor interaction domains in nuclear proteins in which the method comprises analysis of the sequence of a nuclear protein for the presence of signature motifs as defined in any one of claims 1-6 in order to identify an interaction domain or a potential interaction domain.

Fig.1A.

			AAD-ER		ADD-ER Mut	
		LIGAND (E2)	-	100nM	-	100nM
DBD			0.4	0.5	0.3	0.6
DBD-RIP140	20-29	-Y L E G L L M H Q A	2	83	3	2
	132-139	-L L A S L L Q S	3	583	2	3
	184-192	-H L K T L L K K S	0.4	34	0.9	0.6
	266-273	-Q L A L L L S S	0.6	208	0.9	0.9
	379-387	-L L L H L L K S Q	0.5	82	0.7	0.8
	498-506	-V T L L Q L L L G	9	1605	10	11
	712-719	-V L Q L L L G N	2	598	1	2
	818-825	-L L S R L L R Q	0.7	715	1	1
	935-944	-V L K Q L L L S E N	0.7	498	0.9	0.8
	935-942	-V L K Q L L L S	0.2	77	0.4	0.4
DBD-SRC1a	45-53	-E L A E L L S A N	0.5	3.3	0.9	0.6
	111-118	-S L G P L L L E	0.5	0.6	0.6	0.8
	632-640	-K L V Q L L T T T	0.5	678	0.7	0.8
	689-696	-I L H R L L Q E	1	1632	1	0.8
	748-755	-L L R Y L L D K	3	366	3	4
	912-920	-Q L D E L L C P P	0.4	0.7	0.7	0.8
	1434-1441	-L L Q Q L L T E	3	373	3	3
DBD-CBP	68-75	-Q L S E L L R G	0.3	28	0.4	0.5

Fig.1B.

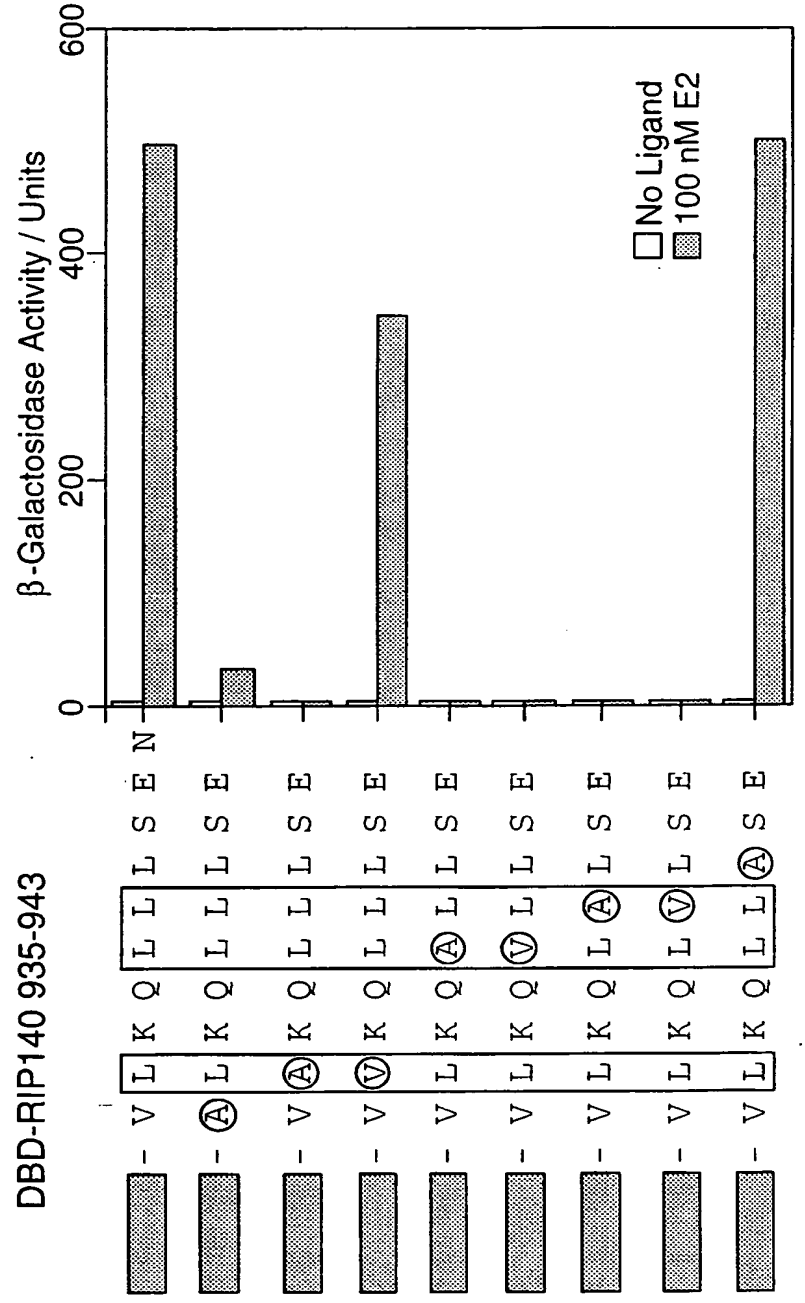

β-Galactosidase Activity / Units
0
200
400
600
No Ligand
100 nM E2
DBD-RIP140 935-943
- V L K Q L L L S E N
- A L K Q L L L S E
- V A K Q L L L S E
- V V K Q L L L S E
- V L K Q A L L S E
- V L K Q V L L S E
- V L K Q L A L S E
- V L K Q L V L S E
- V L K Q L L A S E

Fig.2A.

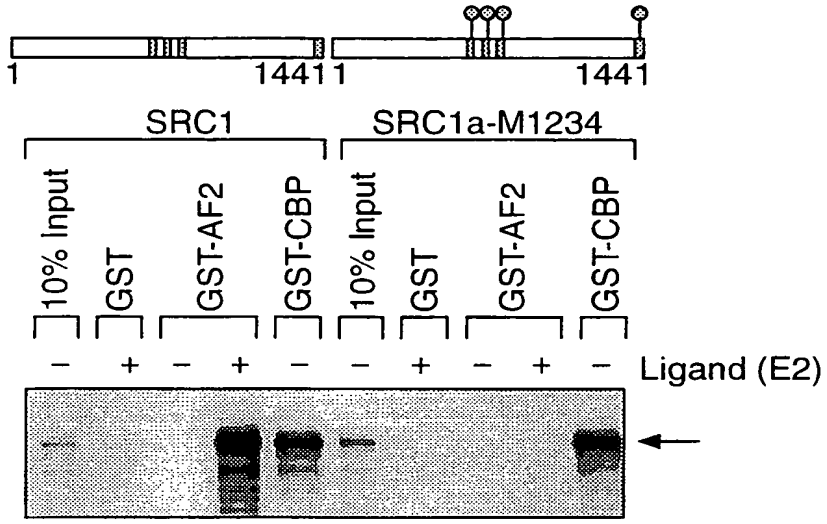
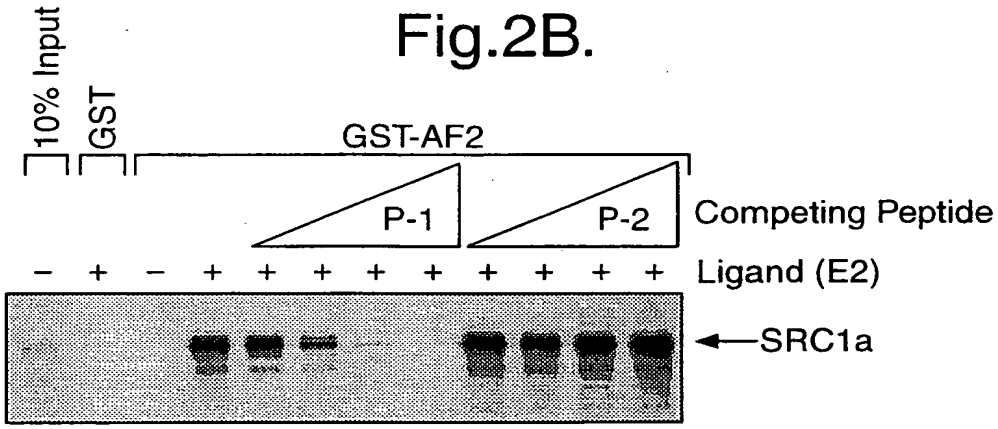


Fig.2B.



P-1 P Q A Q Q K S L L Q Q L L T

P-2 P Q A Q Q K S L L Q Q A A T

Fig.2C.

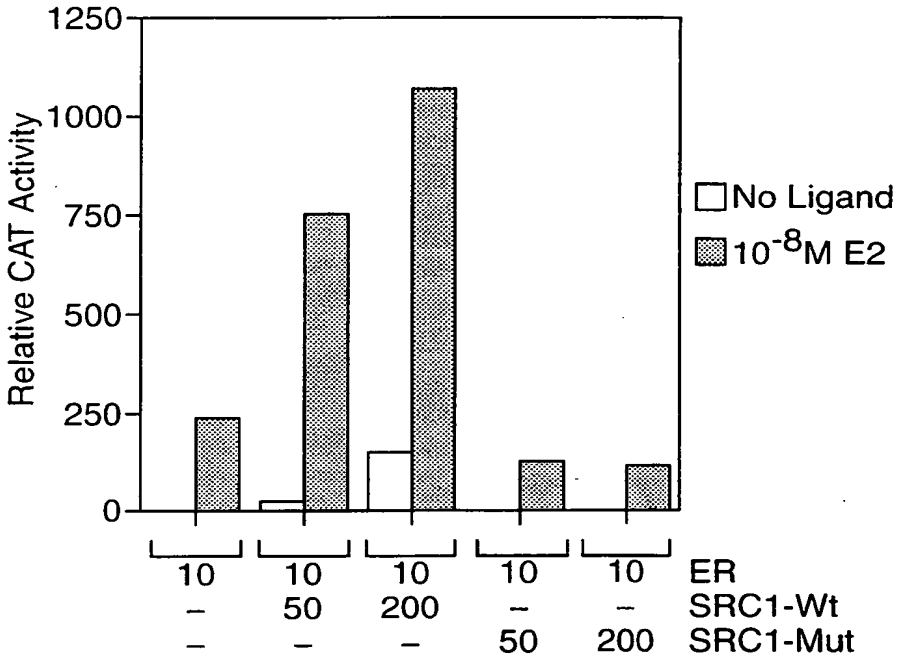

1250
1000
750
500
250
0
Relative CAT Activity
No Ligand
10-8M E2
10 10 10 10 10 ER
– 50 200 – – SRC1-Wt
– – – 50 200 SRC1-Mut

Fig.3A.

		*			*	*						
	Y	L	E	G	L	L	M	H	Q	A	A	20-30
	L	L	A	S	L	L	Q	S	E	S	S	123-142
	H	L	K	T	L	L	K	K	S	K	V	184-194
RIP140	Q	L	A	L	L	L	S	S	E	A	H	266-276
	L	L	L	H	L	L	K	S	Q	T	I	379-389
	L	L	Q	L	L	L	G	H	K	N	E	499-509
	V	L	Q	L	L	L	G	N	P	K	G	712-722
	L	L	S	R	L	L	R	Q	N	Q	D	818-828
	V	L	K	Q	L	L	L	S	E	N	C	935-945
	K	L	V	Q	L	L	T	T	T	A	E	632-642
SRC1a	I	L	H	R	L	L	Q	E	G	S	P	689-699
	L	L	R	Y	L	L	D	K	D	E	K	748-758
	L	L	Q	Q	L	L	T	E				1434-1441
	K	L	L	Q	L	L	T	T	K	S	D	640-650
TIF2	I	L	H	R	L	L	Q	D	S	S	S	689-699
	L	L	R	Y	L	L	D	K	D	D	T	744-754
	Q	L	S	E	L	L	R	G	G	S	G	68-78
CBP	Q	L	V	L	L	L	H	A	H	K	C	356-366
	Q	L	S	E	L	L	R	G	S	S	P	80-90
p300	Q	L	V	L	L	L	H	A	H	K	C	341-351
TIF1	I	L	T	S	L	L	L	N	S	S	Q	722-732
Trip2	M	L	M	N	L	L	K	D	N	P	A	23-33
Trip3	T	L	R	S	L	L	L	N	P	H	L	97-107
Trip4	R	L	A	V	L	L	P	G	R	H	P	36-46
Trip5	E	L	H	N	L	L	E	V	V	S	Q	26-36
Trip8	T	L	R	D	L	L	T	T	T	A	G	36-46
	F	L	D	F	L	L	G	F	S	A	G	73-83
Trip9	V	L	E	L	L	L	R	A	G	A	N	256-266
	I	L	A	R	L	L	R	A	H	G	A	288-298

Fig.3B.

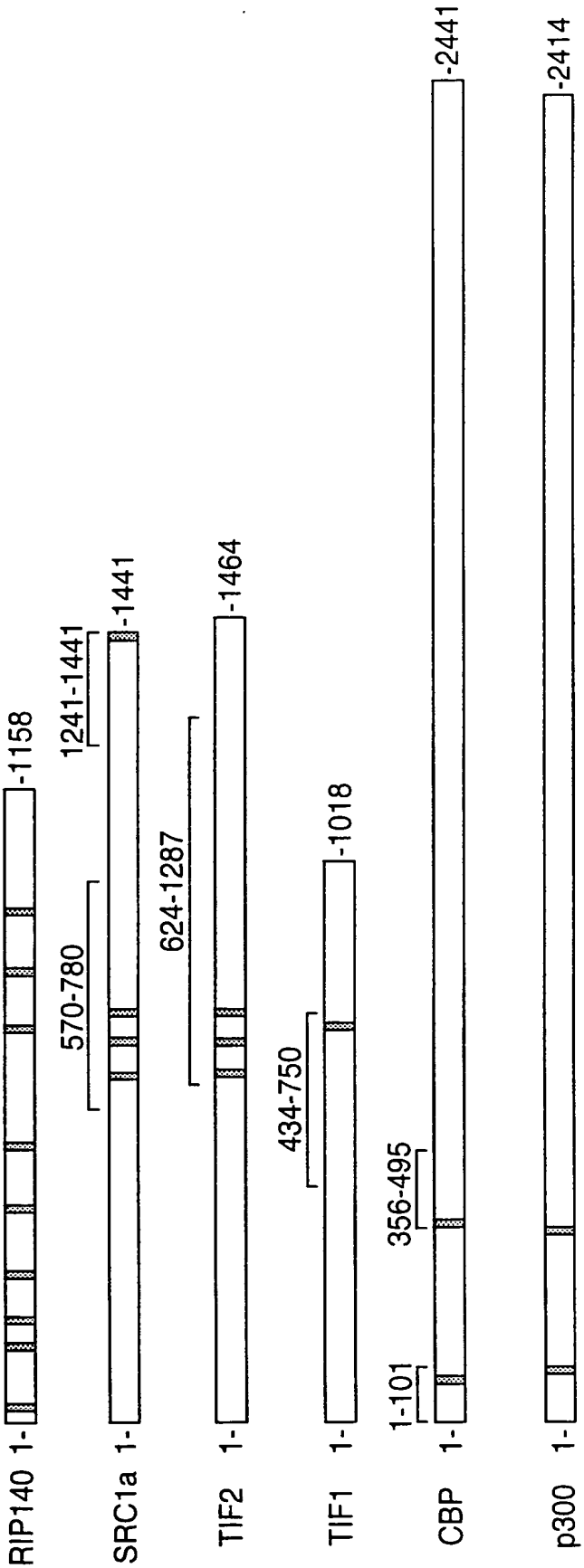
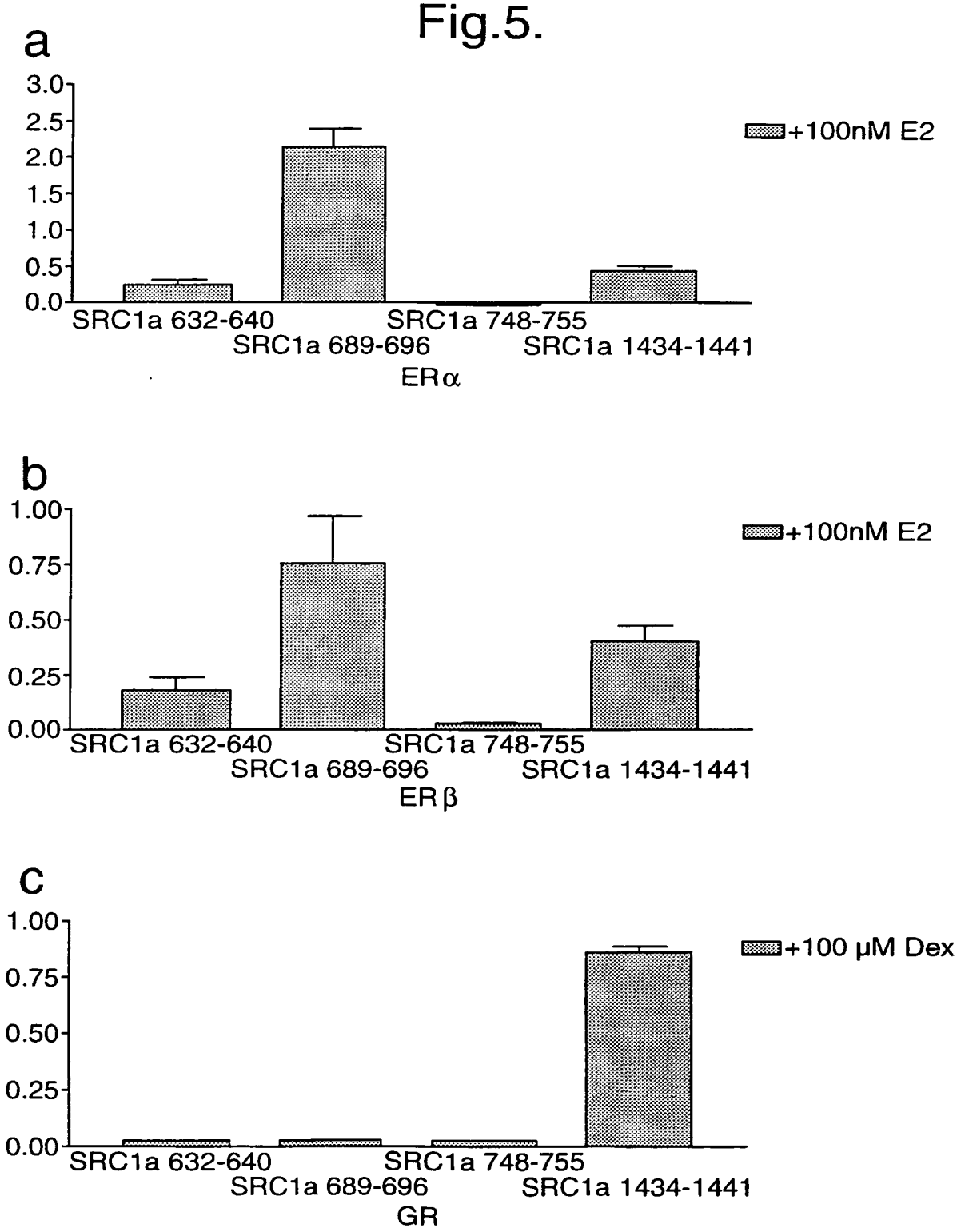

RIP140 1-
27-241
753-981
-1158
SRC1a 1-
570-780
1241-1441
-1441
TIF2 1-
624-1287
-1464
TIF1 1-
434-750
-1018
CBP 1-
1-101
356-495
-2441
p300 1-
-2414

Fig.4.

CBP	A	L	Q	D	L	L	R	T	L	K	S	2067-2077
p300	A	L	Q	N	L	L	R	T	L	R	S	2050-2060
p/CIP	K	L	L	Q	L	L	T	C	S	S	D	613-623
	I	L	H	K	L	L	Q	N	G	N	S	677-687
	L	L	R	Y	L	L	D	R	D	D	P	729-739
ARA70	Q	L	Y	S	L	L	G	Q	F	N	C	91-101
TRIP230	E	L	E	N	L	L	Q	Q	G	G		267-276
	V	L	Q	K	L	L	K	E	K	D		1404-1413
	E	L	N	Q	L	L	N	A	V	K		1524-1533
	V	L	K	D	L	L	K	Q				1971-1978

Fig.5.


a
3.0
2.5
2.0
1.5
1.0
0.5
0.0
+100nM E2
SRC1a 632-640
SRC1a 689-696
SRC1a 748-755
SRC1a 1434-1441
ERα
b
1.00
0.75
0.50
0.25
0.00
+100nM E2
SRC1a 632-640
SRC1a 689-696
SRC1a 748-755
SRC1a 1434-1441
ERβ
c
1.00
0.75
0.50
0.25
0.00
+100 μM Dex
SRC1a 632-640
SRC1a 689-696
SRC1a 748-755
SRC1a 1434-1441
GR

INTERNATIONAL SEARCH REPORT

International Application No
PCT/GB 98/01238

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 G01N33/68 C07K7/06 C07K7/08

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 G01N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	WO 98 11907 A (SQUIBB BRISTOL MYERS CO) 26 March 1998 see claims 1,7,14,15,21 see page 13, line 1 - line 12 see page 15, line 5 - page 25	1-19
P,X	WO 98 02455 A (INST NAT SANTE RECH MED ;UNIV PASTEUR (FR); CENTRE NAT RECH SCIENT) 22 January 1998 see claims 14-19 see page 34, line 4 - page 43, line 2	1-19
P,X	WO 97 45737 A (AMERICAN CYANAMID CO) 4 December 1997 see claims 1-16 see page 4, line 19 - line 26 see page 8, line 18 - page 9, line 7	1-19

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Date of the actual completion of the international search	Date of mailing of the international search report
12 August 1998	20/08/1998

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Routledge, B

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/GB 98/01238

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9811907 A	26-03-1998	AU 4345597 A	14-04-1998
WO 9802455 A	22-01-1998	NONE	
WO 9745737 A	04-12-1997	NONE	

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